

BeneView T5



Patient Monitor

Service Manual

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Revision History

This manual has a revision number. This revision number changes whenever the manual is updated due to software or technical specification change. Contents of this manual are subject to change without prior notice.

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Preface

Manual Purpose

This manual provides detailed information about the assembling, disassembling, testing and troubleshooting of the equipment to support effective troubleshooting and repair. It is not intended to be a comprehensive, in-depth explanation of the product architecture or technical implementation. Observance of the manual is a prerequisite for proper equipment maintenance and prevents equipment damage and personnel injury.

This manual is based on the maximum configuration; Therefore, some contents may not apply to your monitor. If you have any question, please contact our Customer Service Department.

Intended Audience

This manual is for biomedical engineers, authorized technicians or service representatives responsible for troubleshooting, repairing and maintaining the patient monitors.

Abbreviations

Abbreviations used in this manual are:

MPM	multi-parameter module
SMR	satellite module rack
CMS	central monitoring system
PCB	printed circuit board

Passwords

A password may be required to access different modes within the monitor. The passwords are listed below:

- User maintenance: 888888 (User adjustable)
- Factory maintenance: 332888
- Demo mode: 2088
- Configuration mode: 315666 (User adjustable)

FOR YOUR NOTES

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FOR YOUR NOTES

1 Safety

1.1 Safety Information

DANGER

- Indicates an imminent hazard that, if not avoided, will result in death or serious injury.
-

WARNING

- Indicates a potential hazard or unsafe practice that, if not avoided, could result in death or serious injury.
-

CAUTION

- Indicates a potential hazard or unsafe practice that, if not avoided, could result in minor personal injury or product/property damage.
-

NOTE

- Provides application tips or other useful information to ensure that you get the most from your product.
-

1.1.1 DANGER

There are no dangers that refer to the product in general. Specific “Danger” statements may be given in the respective sections of this manual.

1.1.2 Warnings

WARNING

- All installation operations, expansions, changes, modifications and repairs of this product are conducted by authorized personnel.
 - There is high voltage inside the equipment. Never disassemble the equipment before it is disconnected from the AC power source.
 - When you disassemble/reassemble a parameter module, a patient leakage current test must be performed before it is used again for monitoring.
 - The equipment must be connected to a properly installed power outlet with protective earth contacts only. If the installation does not provide for a protective earth conductor, disconnect it from the power line and operate it on battery power, if possible.
 - Dispose of the package material, observing the applicable waste control regulations and keeping it out of children’s reach.
-

1.1.3 Cautions

CAUTION








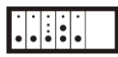













- Make sure that no electromagnetic radiation interferes with the performance of the equipment when preparing to carry out performance tests. Mobile phone, X-ray equipment or MRI devices are a possible source of interference as they may emit higher levels of electromagnetic radiation.
 - Before connecting the equipment to the power line, check that the voltage and frequency ratings of the power line are the same as those indicated on the equipment’s label or in this manual.
 - Protect the equipment from damage caused by drop, impact, strong vibration or other mechanical force during servicing.
-

1.1.4 Notes

NOTE

- Refer to Operation Manual for detailed operation and other information.
-

1.2 Equipment Symbols

	Attention: Consult accompanying documents (this manual).		CIS connector
	Danger: High-voltage		Network connector
	Alternating current(AC)		Defibrillator connector
	Power ON/OFF		Connector for satellite module rack
	Battery indication		Video output
	Zero key		Auxiliary output connector
	Calibrate key		USB connector
	Measure/Standby		Equipotential terminal
	Check sensor		CE marking
	ESD warning symbol for Electrostatic sensitive devices.		
	Type CF applied part. Defibrillator-proof protection against electric shock.		
	Type BF applied part. Defibrillator-proof protection against electric shock.		

FOR YOUR NOTES

2 Theory of Operation

2.1 Introduction

This patient monitor is designed to monitor a fixed set of physiological parameters including ECG, heart rate (HR), respiration (Resp), temperature (Temp), SpO₂, pulse rate (PR), non-invasive blood pressure (NIBP), invasive blood pressure (IBP), cardiac output (C.O.), carbon dioxide (CO₂), oxygen (O₂), anesthetic gas (AG), impedance cardiograph (ICG), bispectral index (BIS), respiration mechanics (RM), continuous cardiac output (PiCCO), central venous oxygen saturation (ScvO₂), electroencephalograph (EEG), and neuromuscular transmission (NMT).

The patient monitor also:

- Provides audible and visual alarm indications in case of patient or equipment problems.
- Enables displaying, reviewing, storing and transferring of real-time data.
- Incorporates multiple input devices such as buttons, knob, touchscreen, keyboard and mouse.
- Interfaces a clinical information system or central monitoring system.
- Enables program upgrade over the network.
- Integrates the information of other devices, which include but are not restricted to anesthesia machine and ventilator.

2.2 System Connections

2.2.1 Mounting the Patient Monitor

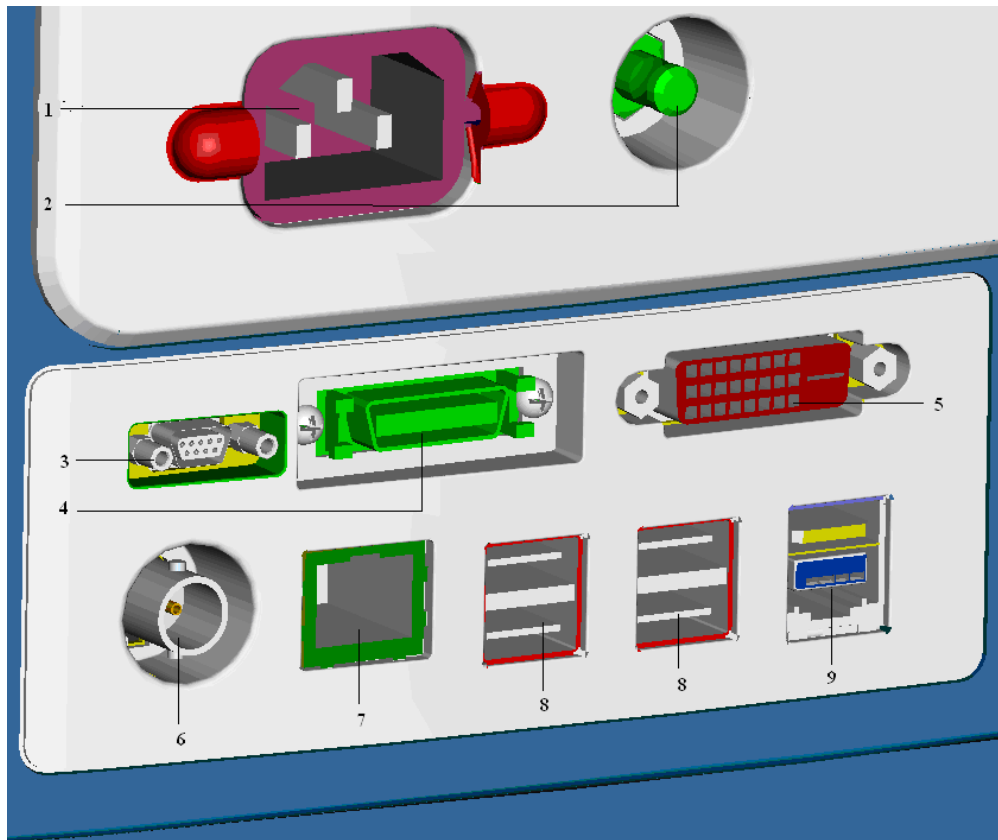
The patient monitor can be mounted on a wall bracket or on a trolley support. The wall bracket or trolley support can be ordered optionally. Each type of mounting bracket is delivered with a complete set of mounting hardware and instructions. Refer to the documentation delivered with the mounting hardware for instructions on assembling mounts.

CAUTION

- **Not using screw and bracket specified by Mindray may cause the screw to touch the internal battery and lead to monitor damage (the proper screw depth should be between 6.5mm and 7.5mm). Take out the rubber stoppers in the two screw holes of the base when installing the tray.**
 - **The mounting bracket should be installed by our qualified service personnel, or engineers who have adequate knowledge on it.**
 - **If other mounting solution is used, the installation personnel and the customer should verify if it can be safely used on the patient monitor, and the customer assume the responsibility for any risk resulting from that.**
-

2.2.2 Connectors for Peripheral Devices

On the back of the patient monitor you will find all connectors for peripheral devices.



1. AC Power Connector: used to connect an AC power source (100 to 240 VAC, 50/60Hz).
2. Equipotential Terminal: used to connect the equipotential terminal of other equipment, eliminating potential difference between different pieces of equipment.
3. Analog Output and Defibrillator Connector: It is a Micro-D connector used to output analog signals and defibrillator synchronization signals.
4. CIS Connector: It is used to connect a CIS.
5. Video Output: It is a DVI-D connector used to connect a secondary display.
6. Aux Output Connector: It is a BNC connector used to output nurse call signals.
7. Network Connector: It is a RJ45 connector used to connect an ethernet network or a PC.
8. USB Connector: used to connect any USB-compatible peripheral device.
9. SMR Connector: It is used to connect the SMR and outputs a 12V DC.

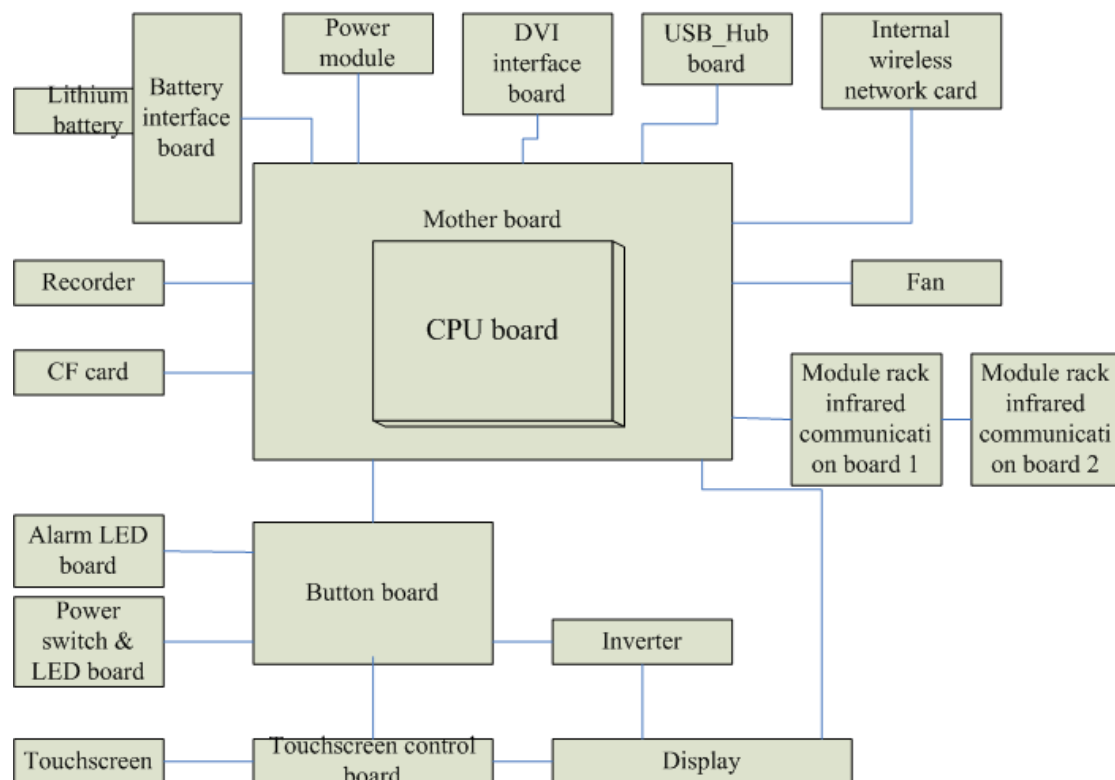
2.3 Main Unit

The patient monitor consists of:

- Input system: button board, knob, touchscreen, power switch and LED board
- Output system: LCD panel, alarm LED board, recorder and speaker
- Processing and communications system: main board and integral module rack assembly.
- Power management system: battery, battery interface board and power module
- Equipment interface system: USB_Hub interface board, DVI interface board CF card assembly and internal wireless network card.

Additionally, the patient monitor can also connect satellite module rack (SMR), parameter modules, BeneLink module, mouse, keyboard, etc.

The following diagram illustrates the structure of the patient monitor



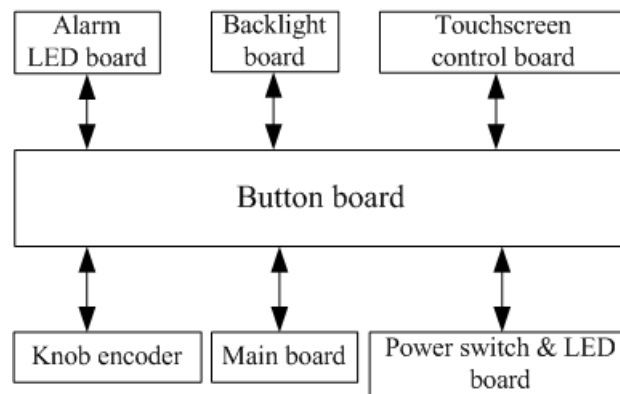
2.3.1 Input System

Button board

The button board, located at the lower part of the monitor's front panel, contains 6 keys and provides connections for the following components to the main board:

- Knob
- Power switch & LED board
- Touchscreen control board
- Alarm LED board
- Inverter

The following diagram shows the button board connections.



Knob

The knob can be pressed, or rotated both clockwise and counter-clockwise. It is connected with the button board.

Touchscreen

The touchscreen enables touch operations and can be calibrated. It is connected with the touchscreen control board and main board.

Power switch & LED Board

The power switch & LED board controls the power supply for the main unit. It has three LEDs, which respectively indicate the AC power status, battery status and monitor power on/off status. It is connected with the button board.

2.3.2 Output System

LCD

The patient monitor adopts a high-resolution LCD. The LCD is connected with the main board. The signals and power supply from the backlight board are transferred by the button board.

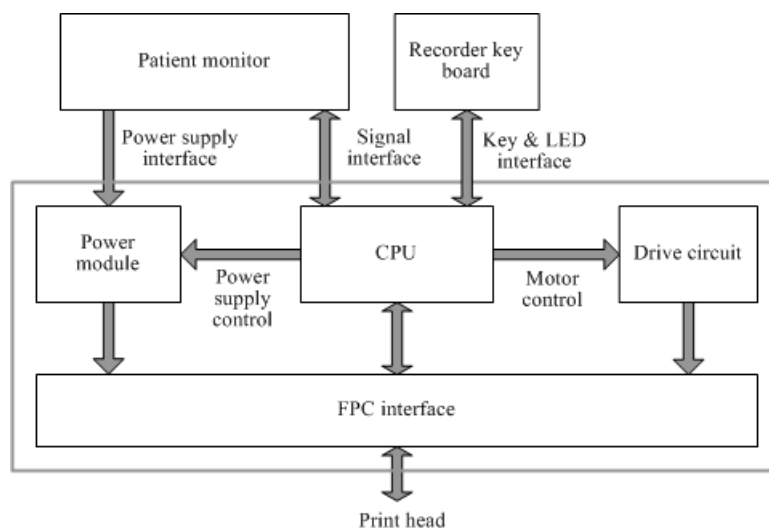
Alarm Lamp

The patient monitor has two alarm lamps: alarm lamp and technical alarm lamp. Alarm lamp lights either red or yellow whereas technical alarm lamp lights blue only. The alarm lamp signals are transferred by the button board and are directly controlled by the main board

Recorder

The recorder receives data form the main board and then sends them to the thermal printhead for printing. The recorder has a hardkey (starting/stopping recordings) and a green LED (It is ON during normal working) on its front panel. It is connected with the mother board.

The following diagram shows its operating principle.



Module	Description
Power interface	Introduces a DC from the main board.
Recorder power module	Converts the input power into voltages that fit each module and then forwards them to each module.
Recorder CPU	Controls the communications between modules.
Signal interface	Controls the communications between the main board and the recorder CPU.
Motor drive circuit	Receives the control signals from the CPU and then forwards them to the step motors
Button & LED board	Includes one button and one LED which are directly controlled by the CPU.

Speaker

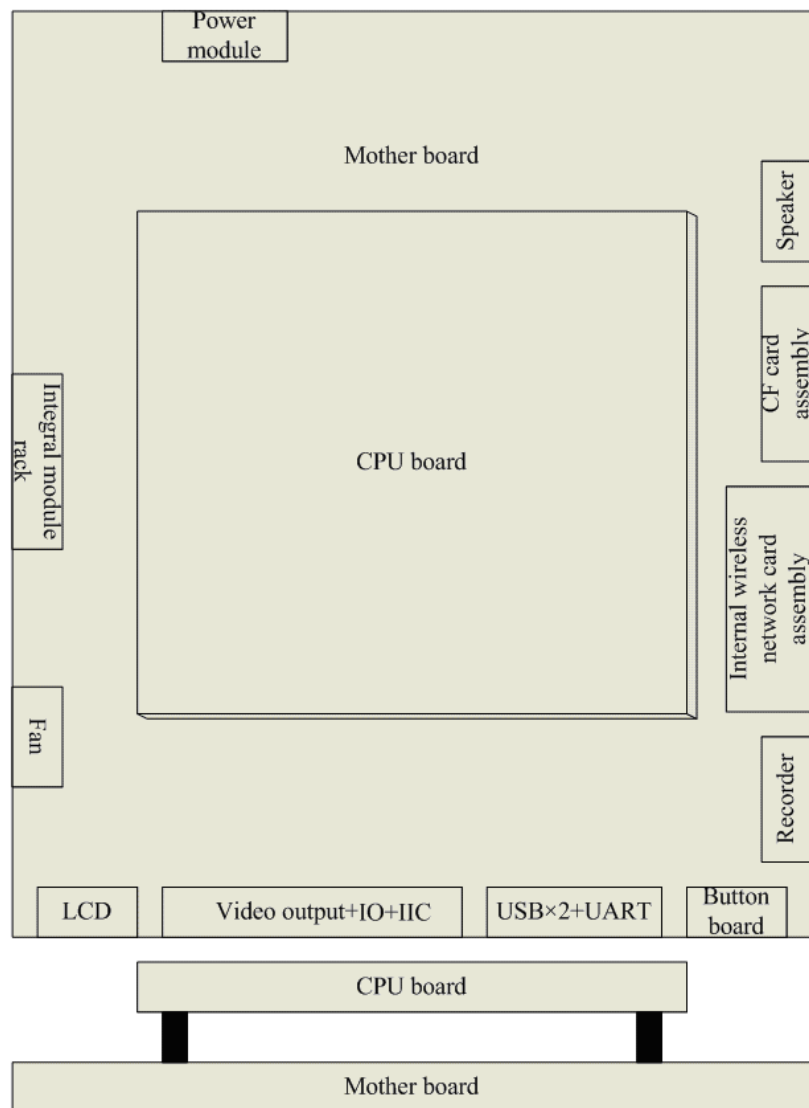
The speaker provides sound for alarms, key strokes, heart beats and pulse, and allows PITCH TONE and multi-level tone modulation. It is connected with the main board and is directly driven by the main board.

2.3.3 Processing and Communications System

Main Board

The main board is the heart of the patient monitor. It implements a series of tasks including input & output control, data storage and processing, display processing, system control, communication management, printing management and alarming, etc.

The main board comprises the CPU board and mother board. The following diagram shows interfaces to other components.



The CPU board is an essential CPU system containing the CPU, FLASH, memory, realtime clock, EEPROM, etc. It interfaces to the mother board only, which then provides interfaces to all other external devices.

The mother board is in charge of connections and communications with other components and provides the following interfaces:

Name	Description
LCD connector	Connects the built-in display.
Video output +IO +IIC	Connects the digital video interface board.
USB×2+network+RS422 +GPIO port	Connects the USB_Hub board.
Button board connector	Connects the button board.
Recorder connector	Connects the recorder.
CF card connector	Connects the CF card assembly.
Speaker connector	Connects the speaker.
Power module connector	Connects the power module.
Integral module rack connector	Connects the 3-slot rack communication board in the integral module rack.
Fan connector	Connects the fan.
Internal wireless network card assembly	Connects the internal wireless network card.

Integral Module Rack

The patient monitor has two kinds of integral module rack: 2-slot and 5-slot. The control board includes a NIOS II FPGA. It implements protocol conversion and infrared communication between the main unit and the parameter modules

The module rack communication board can be a 2-slot type or a 3-slot type. The 3-slot communication board communicates the main board directly. The 2-slot communication board is connected with and controlled by the 3-slot communication board. The 3-slot communication board has the function of communication control. The 2-slot communication board consists of the infrared circuit and module power circuit. The RS422 drive circuit is located on the 3-slot communication board.

2.3.4 Power Management System

Battery

The patient monitor uses two chargeable lithium-ion batteries (11.1 V, 4500 mAh). The battery compartment is located at the bottom of the patient monitor. The battery power is supplied to the mother board via the battery interface board, and then to the power module.

Battery Interface Board

The Battery interface board connects the batteries to the DC input terminal of the power module via the mother board, implementing charging and discharging of the batteries and the power board.

Power Module

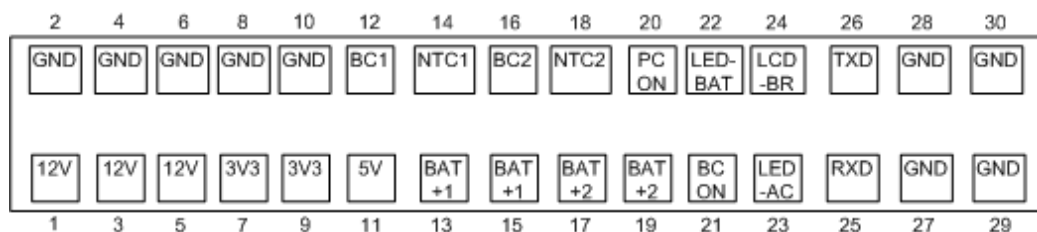
The power module is located at the back of the patient monitor. The main part of the power module is the power board, which contains charging & power management board, voltage drop DC transforming board and voltage rise and drop DC transforming board.

The power module converts the input power into DC power supplies and then distributes them to each component of the patient monitor. The input power comes from either the batteries or an AC source. The patient monitor will run power from the AC source whenever an AC source is available. If the AC source becomes unavailable, the patient monitor will automatically switch to the battery power. This does not affect the monitor's operating status.

The power module protects itself and the patient monitor by switching off AC input or DC output in case of overcurrent, short circuit and overvoltage. The power module provides 3 DC outputs:

Outputs	Description
+3.3 V	Power supply of the LCD, mother board, CPU board, DVI interface board and integral module rack.
+5.0 V	Power supply of the DVI interface board, recorder, CF storage card board and USB_Hub board.
+12 V	Power supply of the recorder, LCD inverter, integral module rack, parameter modules, USB_Hub board.

The following diagram shows the pins of the power socket connecting the power module and the mother board:

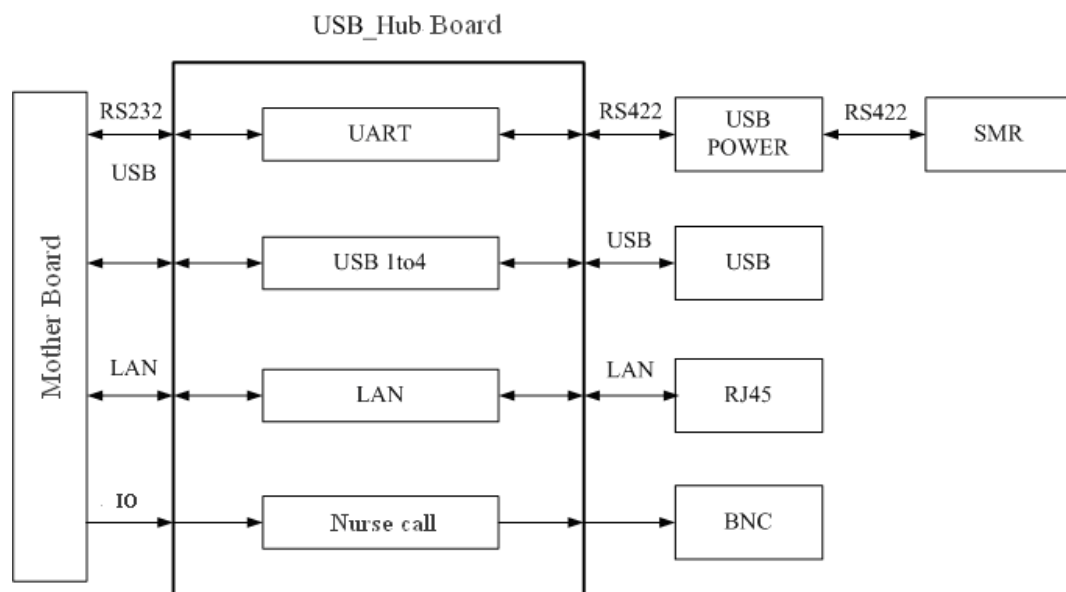


Pin ID	Marking	Description
1/3/5	12V	The positive output of the 12 VDC power
2/4/6/8/10/ 27/28/29/30	GND	The output grounding terminal of the power board.
7/9	3V3	The positive output of the 3.3 VDC power
11	5V	The positive output of the 5 VDC power
12	BC1	Signal indicating whether battery 1 is available. Low level indicates that battery 1 is available and high level indicates that battery 1 is not available.
13/15	BAT+1	Input of battery 1, connecting to the positive pole of the battery.
14	NTC1	Thermistor signal of battery 1.
16	BC2	Signal indicating whether battery 2 is available. Low level indicates that battery 2 is available and high level indicates that battery 2 is not available.
18	NTC2	Thermistor signal of battery 2.
17/19	BAT+2	Input of battery 2, connecting to the positive pole of the battery.
20	PCON	Power on/off control signal. It is a TTL pulse signal inputted from the back board. Every time when the power on/off switch is pressed (pulse of falling edge), a switch between power “on” and “off” happens. The pulse duration is no less than 0.1 s for power on, 2 s for power off and 10 s for illegal power off.
21	BCON	Backlight on/off signal and switch output signal. The main board sends the LCD backlight on/off signals to the power board via a serial port, the power board processes the signals and output them. Low level is output when the backlight is off and high level is output when the backlight is on.
22	LED-BAT	Battery status indication driving output
23	LED-AC	AC power status indication signal
24	LCD-BR	Backlight brightness control voltage.

2.3.5 Equipment Interface System

USB_Hub board

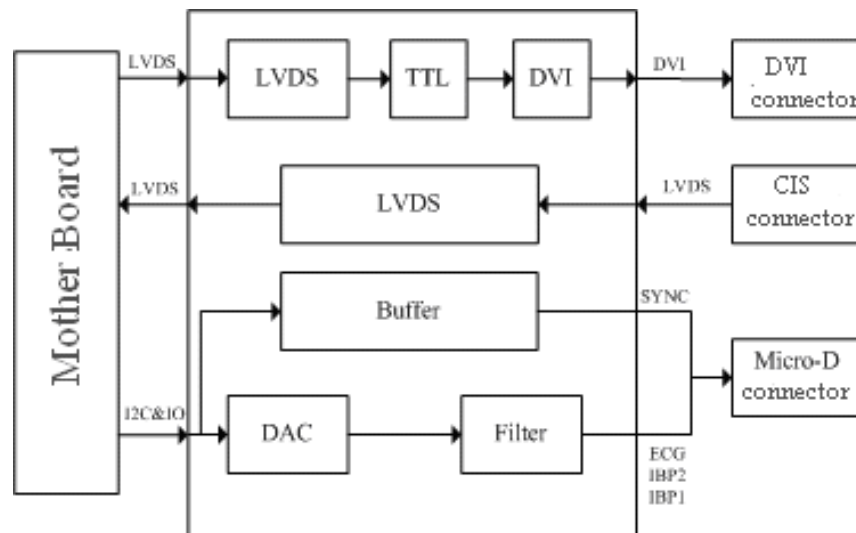
The USB_Hub board is connected with the mother board. It is compatible with USB1.1 connectors and supports equipment hot plug. The UART signal output by the main board is converted into RS422 signal by the USB_HUB board. It receives 5 VDC and 12 VDC inputs from the power module, of which the 5 VDC is supplied to the USB interface board and the 12 VDC is outputted to the SMR connector through a fuse.



BNC	It is a BNC connector used to output nurse call signals.
RJ 45 connector	It is a standard RJ45 connector, providing 10/100 BASE-TX Ethernet communications channels. It connects an Ethernet network or a PC.
USB connector	Connects devices with USB connector.
USB&POWER connector	Provides RS232 and RS422 interfaces for the communication between main board and SMR. It receives 5 VDC and 12 VDC inputs from the power module, of which the 5 VDC is supplied to the USB interface board and the 12 VDC is outputted to the SMR connector through a fuse.

DVI Interface Board

The DVI interface board is connected with the mother board. The following diagram shows its interfaces to other components.



Interface	Description
DVI connector	Connects the secondary display.
CIS Connector	Connects the CIS system.
Micro-D connector	Outputs analog signals and defibrillator synchronization signals.

CF Card Assembly

The CF assembly serves the non-volatile CF card which is used for data storage and transferring. It is connected with the mother board.

Internal wireless network card

The internal wireless network card connects with the mother board. User can set network type as LAN or WLAN through user interface and can set the internal wireless network card through PC.

2.4 Parameter Module

Each parameter module may consist of the module infrared communication board, module power board, module button board, parameter board, etc.

2.4.1 Module Infrared Communication Board

The module infrared communication board allows a short delay when powering up the module and adopts FPGA to enable infrared communications between the module and the module rack. An ID is integrated into the module infrared communication board. When a module is inserted in the module rack, the ID is automatically sent to the module rack.

2.4.2 Module Power Board

Some modules have no power board. There are two kinds of module power board:

1. Isolated power board: converts the 12 V DC into a 12 V isolated DC and a 5 V isolated DC.
2. Non-isolated power board: converts the 12 V DC into a 5 V DC

2.4.3 Module Button Board

There are keys and a LED on the module button board.

2.4.4 Parameter Board

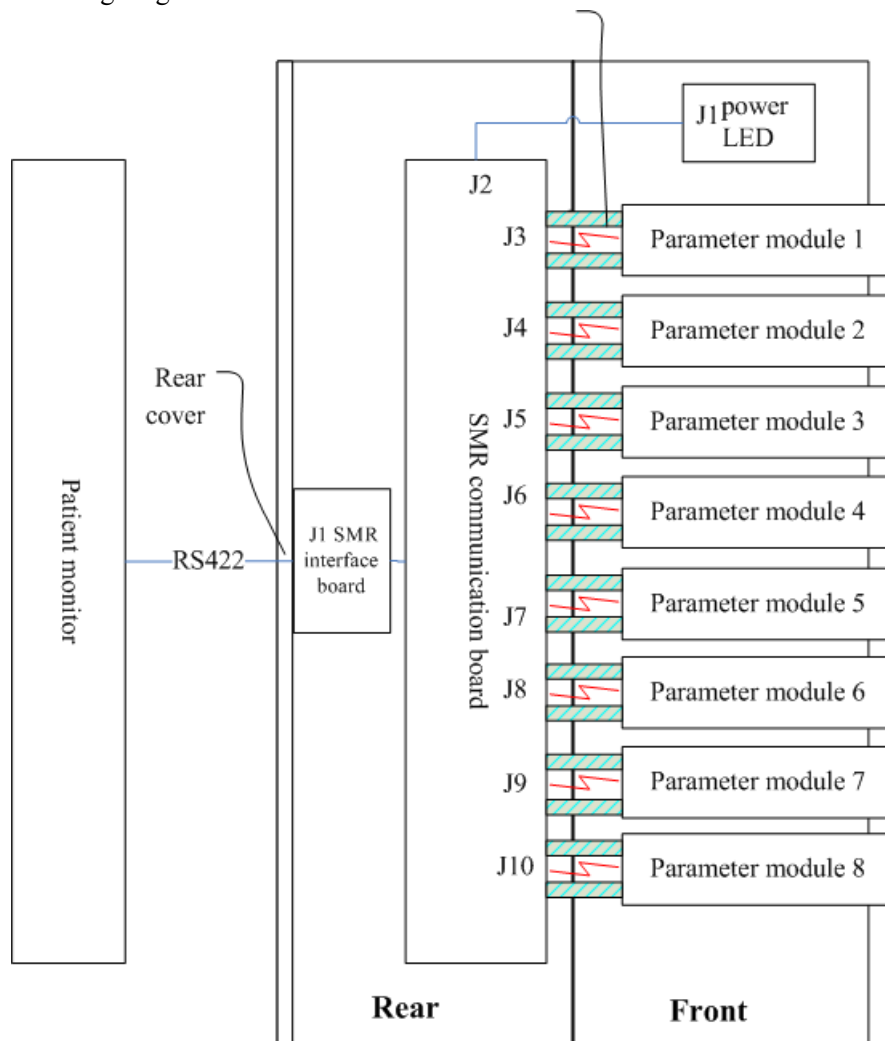
The parameter board is a parameter measurement component, which is the most important component of the parameter module.

2.5 SMR

The satellite module rack (SMR) is independent of the patient monitor. It provides 8 slots for mounting parameter modules. It has the following features:

- It allows a parameter module to be plugged and unplugged with the patient monitor on. This allows function extension and patient transfer.
- It does not have its own power supply. It is run by 12 V DC supplied by the patient monitor and then supplies power supply to each parameter module via the contact screws.
- It implements communication protocol conversions between the patient monitor and each parameter module, provides infrared communications for parameter modules, and is responsible for detecting infrared communications malfunction for each parameter module.

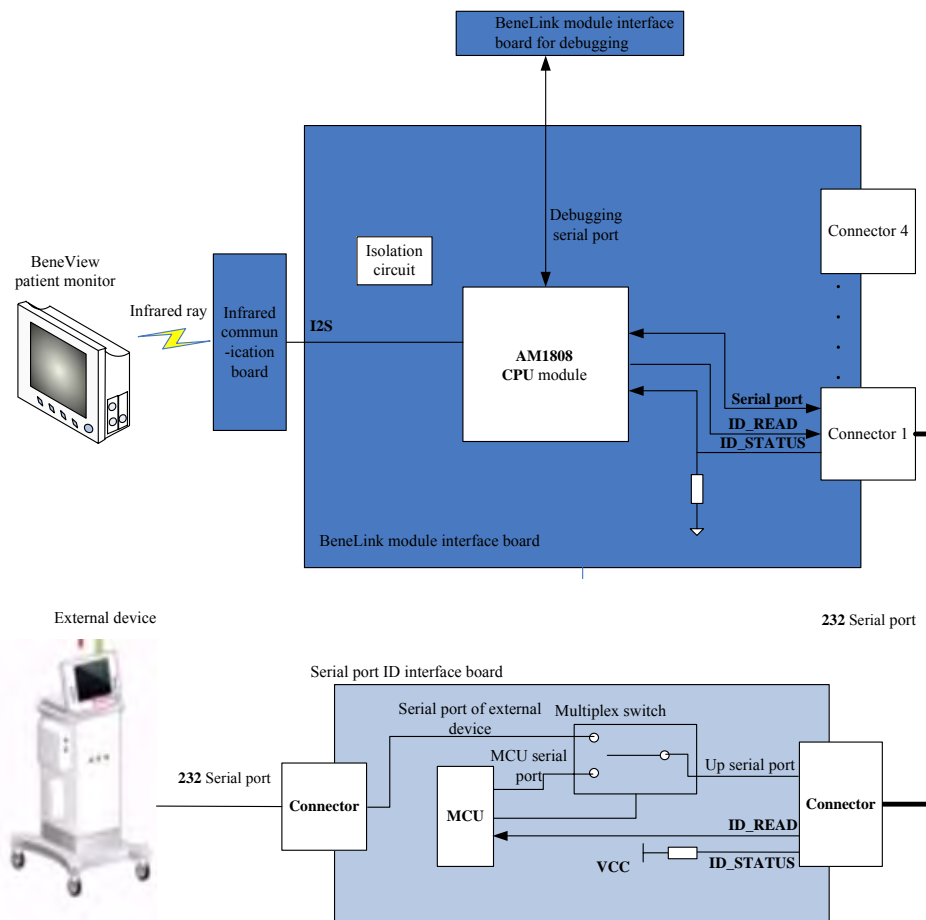
The following diagram shows the structure of the SMR.



2.6 BeneLink Module

BeneLink module allows the information (patient data, alarms, etc.) from the external device to be displayed, saved, recorded, printed, or calculated through a BeneView patient monitor. If the patient monitor is connected with the CMS or gateway, information from the external device can also be transmitted to the CMS or gateway. BeneLink module connects with the external device through an ID module, which enables the information transmission between the BeneLink module and the external device. BeneLink module can be connected to many external devices such as anesthesia machine and ventilator.

The following diagram shows the structure of the BeneLink module:



3 Testing and Maintenance

3.1 Introduction

To ensure the patient monitor always functions normally, qualified service personnel should perform regular inspection, maintenance and test. This chapter provides a checklist of the testing procedures for the patient monitor with recommended test equipment and frequency. The service personnel should perform the testing and maintenance procedures as required and use appropriate test equipment.

The testing procedures provided in this chapter are intended to verify that the patient monitor meets the performance specifications. If the patient monitor or a module fails to perform as specified in any test, repairs or replacement must be done to correct the problem. If the problem persists, contact our Customer Service Department.

CAUTION

- All tests should be performed by qualified service personnel only.
 - Care should be taken to change the settings in [User Maintenance>>] and [Factory Maintenance>>] menus to avoid loss of data.
 - Service personnel should acquaint themselves with the test tools and make sure that test tools and cables are applicable.
-

3.1.1 Test Equipment

See the following sections.

3.1.2 Test Report

Upon completion of the tests, the table of preventative maintenance test reports and the table of maintenance test reports in this chapter should be kept properly.

3.1.3 Preventative Maintenance

Below are preventative maintenance tests which need to be performed on the monitor. See the following sections for detailed maintenance procedures.

- Visual inspection
- NIBP test and calibration
- Microstream and Sidestream CO₂ test and calibration
- AG test and calibration

3.1.4 Recommended Frequency

Check/Maintenance Item		Frequency
Preventative Maintenance Tests		
Visual inspection		1. When first installed or reinstalled.
NIBP test	Pressure check	1. If the user suspects that the measurement is incorrect. 2. Following any repairs or replacement of relevant module. 3. At least once a year. 4. AG leakage test should be performed before AG measurement.
	Leakage test	
	Calibration	
Sidestream and Microstream CO ₂ tests	Leakage test	
	Performance test	
	Calibration	
AG tests	Performance test	
	Calibration	
Performance Tests		
ECG test	Performance test	1. If the user suspects that the measurement is incorrect. 2. Following any repairs or replacement of relevant module. 3. At least once every two years. At least once a year is recommended for NIBP, CO ₂ , NMT and AG. 4. AG leakage test should be performed before AG measurement.
	Calibration	
Resp performance test		
SpO ₂ test		
NIBP test	Pressure check	
	Leakage test	
	Calibration	
Temp test		
IBP test	Performance test	

	Pressure calibration	
C.O. test		
Mainstream CO ₂ test		
Sidestream and Microstream CO ₂ tests	Leakage test	
	Performance test	
	Calibration	
AG tests	Leakage test	
	Performance test	
	Calibration	
ICG test		
BIS test		
RM test		
CCO/SvO ₂ test	Interconnecting function	
	Output calibration	
NMT test	Performance test	
	Sensor check	
EEG test		
PiCCO test		
ScvO ₂ test		
Nurse call relay performance test		If the user suspects that the nurse call or analog output does not work well.
Analog output performance test		
Electrical Safety Tests		
Electrical safety tests		Refer to <i>A Electrical Safety Inspection</i> .
Other Tests		
Power on test		1. When first installed or reinstalled. 2. Following any maintenance or the replacement of any main unit parts.
Touchscreen calibration		1. When the touchscreen appears abnormal. 2. After the touchscreen is replaced.
Recorder check		Following any repair or replacement of the recorder.
Network print test		1. When first installed. 2. Whenever the printer is serviced or replaced.
Device integration check		1. When first installed. 2. Following any repair or replacement of the external device.
Battery check	Functionality test	1. When first installed. 2. Whenever a battery is replaced.

3.2 Preventative Maintenance Procedures

3.2.1 Visual Inspection

Inspect the equipment for obvious signs of damage. The test is passed if the equipment has no obvious signs of damage. Follow these guidelines when inspecting the equipment:

- Carefully inspect the case, display screen, buttons and knob for obvious signs of damage.
- Inspect the SMR and parameter modules for obvious signs of damage.
- Inspect the power cord, wall-mount bracket and module accessories for obvious signs of damage.
- Inspect all external connections for loose connectors, bent pins or frayed cables.
- Inspect all connectors on the equipment for loose connectors or bent pins.
- Make sure that safety labels and data plates on the equipment are clearly legible.

3.2.2 NIBP Tests

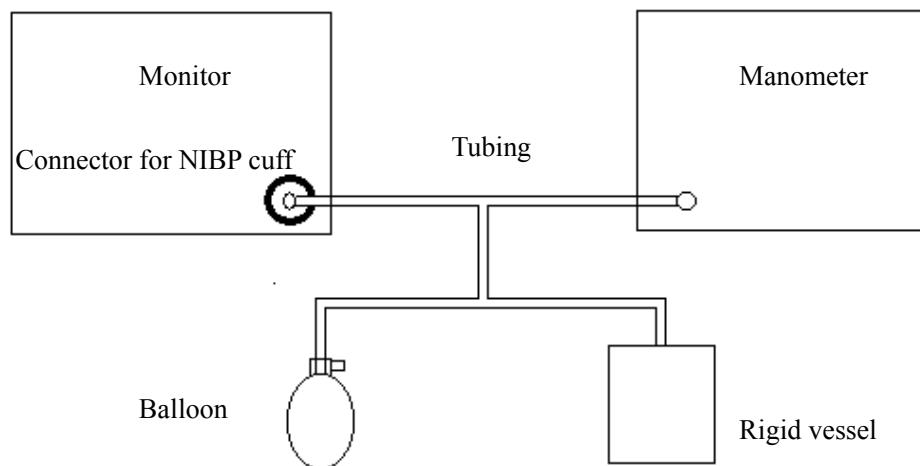
NIBP Accuracy Test

Tools required:

- T-shape connector
- Appropriate tubing
- Balloon pump
- Rigid Vessel with volume 500 ± 25 ml
- Reference manometer (calibrated with accuracy equal to or greater than 1 mmHg)

Follow this procedure to perform the test:

1. Connect the equipment as shown below.



2. Before inflation, the reading of the manometer should be 0. If not, turn off the balloon pump to let the whole airway open to the atmosphere. Turn on the balloon pump after the reading is 0.
3. Select [**Main Menu**]→ [**Maintenance >>**]→ [**NIBP Accuracy Test**].
4. Check the manometer values and the monitor values. Both should be 0mmHg.
5. Raise the pressure in the rigid vessel to 50 mmHg with the balloon pump. Then, wait for 10 seconds until the measured values become stable.
6. Compare the manometer values with the monitor values. The difference should be 3 mmHg. If it is greater than 3 mmHg, contact your service personnel.
7. Raise the pressure in the rigid vessel to 200 mmHg with the balloon pump. Then, wait for 10 seconds until the measured values become stable and repeat step 6.

NOTE

- You can use an NIBP simulator to replace the balloon pump and the reference manometer to perform the test.
 - You can use an appropriate cylinder and a cuff instead of the rigid vessel.
-

NIBP Leakage Test

NOTE

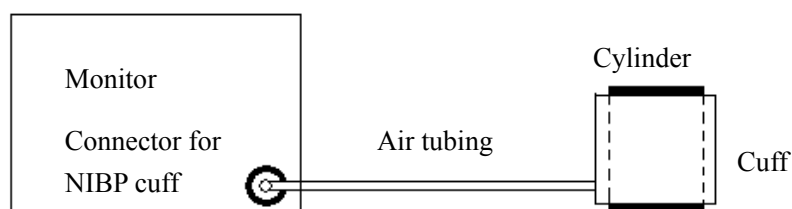
- You should perform NIBP accuracy test and make sure the test result is pass prior to NIBP leakage test.
-

Tools required:

- NIBP cuff for adult patient
- Appropriate tubing
- Cylinder

Follow this procedure to perform the test:

1. Set [**Patient Cat.**] to [**Adu**].
2. Connect the NIBP cuff with the NIBP connector on the monitor.
3. Apply the cuff to the cylinder as shown below.



4. Select [**Main Menu**]→ [**Maintenance>>**]→ [**NIBP Leakage Test**]. The message [**Leakage Testing...**] is displayed in the NIBP parameter area.
5. The cuff automatically deflates after 20s, which means NIBP leakage test is completed. If no message is displayed in the NIBP parameter area, it indicates that the system has no leakage. If the message [**NIBP Pneumatic Leak**] is displayed, it indicates that the system may have a leakage. In this case, check if all connections are good and the cuff and tubing have no leakage. Perform the test again after making sure all connections are good and the cuff and tubing have no leakage.

You can either perform a manual leakage test:

1. Perform procedures 1-4 in the ***NIBP Accuracy Test*** section.
2. Raise the pressure in the rigid vessel to 250 mmHg with the balloon pump. Then, wait for 5 seconds to let the measured values becoming stable.
3. Record the current pressure value and meanwhile use a time counter to count time. Then, record the pressure value after counting to 60s.
4. Compare the two values and make sure the difference should not be greater than 6 mmHg.

3.2.3 Sidestream and Microstream CO₂ Module Tests

Leakage test

Follow this procedure to perform the test:

1. Plug the module into the module rack.
2. Wait until CO₂ warmup is finished and then use your hand or other objects to completely block the gas inlet of the module or watertrap. The sidestream and microstream CO₂ modules will behave as follows:

Sidestream: The alarm message [**CO₂ FilterLine Err**] is displayed on the screen after 3 seconds. Block the gas inlet for another 60 s. Select [**Main Menu**]→ [**Maintenance >>**]→ [**User Maintenance >>**] → enter the required password → [**Maintain CO₂ >>**]→ [**Calibrate CO₂ >>**], and check that the flow rate is less than 10ml/min. The module does not leak if current flow rate is less than 10ml/min and the alarm message does not disappear.

- ◆ Microstream: The alarm message [**CO₂ Purging**] is displayed on the screen after certain time. Block the gas inlet for another 30s. If alarm message [**CO₂ FilterLine Err**] is shown, it indicates that the module does not leak.

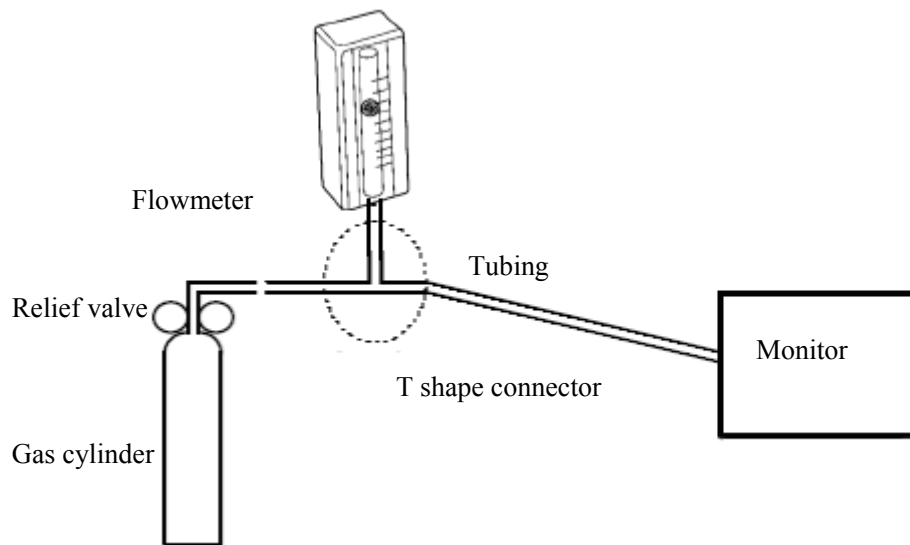
Accuracy Test

Tools required:

- A steel gas cylinder with $6 \pm 0.05\%$ CO₂ and balance gas N₂
- T-shape connector
- Tubing
- Flowmeter

Follow this procedure to perform the test:

1. Plug the module into the module rack.
2. Wait until the CO₂ module warmup is finished, and check the airway for leakage and perform a leakage test as well to make sure the airway has no leakage.
3. Enter [User Maintenance]→ [Maintain CO₂ >>]→ [Calibrate CO₂ >>].
4. Connect the test system as follows:



5. Open the relief valve, and adjust it until the flowmeter has a stable reading between 10 ml/min and 50ml/min.
6. Check the realtime CO₂ value is within $6 \pm 0.05\%$ in the [Calibrate CO₂] menu.

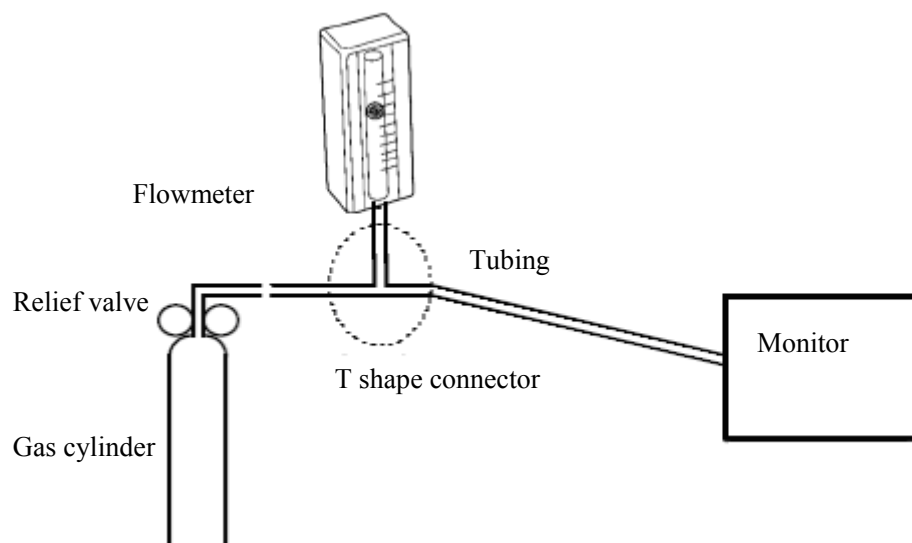
Calibration

Tools required:

- A steel gas cylinder with $6 \pm 0.05\%$ CO₂ and balance gas N₂
- T-shape connector
- Tubing
- Flowmeter

Follow this procedure to perform a calibration:

1. Make sure that the sidestream or microstream CO₂ module has been warmed up or started up.
2. Check the airway for leakage and perform a leakage test as well to make sure the airway has no leakage.
3. Select [**Main Menu**] → [**Maintenance >>**] → [**User Maintenance >>**] → enter the required password → [**Maintain CO₂ >>**] → [**Calibrate CO₂ >>**].
4. In the [**Calibrate CO₂**] menu, select [**Zero**].
5. After the zero calibration is finished successfully, connect the equipment as follows:



6. Open the relief valve, and adjust it until the flowmeter has a stable reading between 10 ml/min and 50ml/min.
7. In the [**Calibrate CO₂**] menu, enter 6% (the CO₂ concentration) in the [**CO₂**] field.
8. In the [**Calibrate CO₂**] menu, the measured CO₂ concentration is displayed. After the measured CO₂ concentration becomes stable, select [**Calibrate CO₂**] to calibrate the CO₂ module.

If the calibration is finished successfully, the message [**Calibration Completed!**] is displayed in the [**Calibrate CO₂**] menu. If the calibration failed, the message [**Calibration Failed!**] is displayed. In this case, perform another calibration.

3.2.4 AG Tests

Leakage Test

Follow this procedure to perform the test:

1. Plug the AG module into the module rack.
2. Wait until the AG module warmup is finished and then use your hand or other objects to completely block the gas inlet of the AG module. An alarm message [**AC Airway Occluded**] will appear on the screen.
3. Block the gas inlet for another 30 s. Select [**Main menu**]→[**Maintenance >>**]→[**User Maintenance >>**]→enter the required password→[**Calibrate AG >>**], and check that current flow rate is less than 10ml/min.

If the alarm message does not disappear, it indicates that the module does not leak.

If the flow rate is less than 10ml/min and the alarm message [**AC Airway Occluded**] does not disappear, it indicates that the module does not leak. If the alarm message disappear, or the flow rate is greater than or equal to 10ml/min, it indicates that the module leaks. If the problem remains, contact your service personnel for help.

Accuracy Test

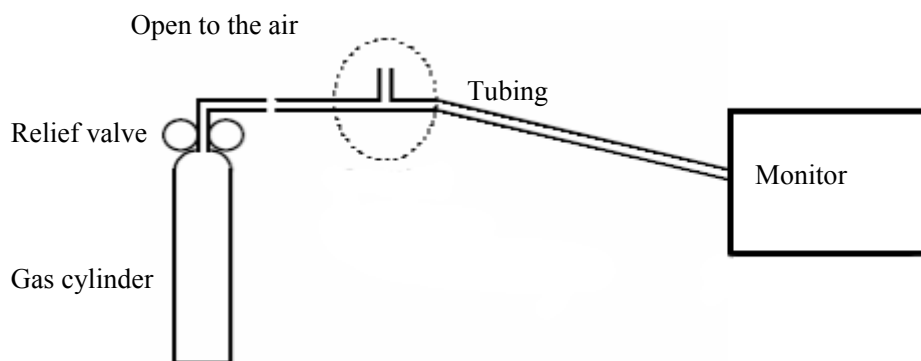
Tools required:

- Gas cylinder with 100% O₂ and a certain standard gas (such as $6 \pm 0.05\%$ CO₂, Bal N₂), or standard gas mixture. Gas concentration should meet the following requirements :
- AA \geq 1.5%, CO₂ \geq 1.5%, N₂O \geq 40%, O₂ \geq 40%, of which AA represents an anesthetic agent. $a/c \leq 0.01$ (a is the gas absolute concentration accuracy; c is the gas concentration)
- T-shape connector
- Tubing

Follow this procedure to perform the test:

1. Plug the AG module into the module rack.
2. Wait at least 10 min and then perform a leakage test to make sure the airway has no leakage.
3. Check if the fan inside the AG module works correctly.

-
4. Connect the test system as follows:



5. Open the relief valve and vent a standard gas and make sure that there is an excess gas flow through the T-shape connector to air.
6. Check that the concentration of each composition meets the specification stated in the Operator's Manual.

WARNING

- When performing AG accuracy test and AG calibration, be sure to dispose of exhaust gas properly.
-

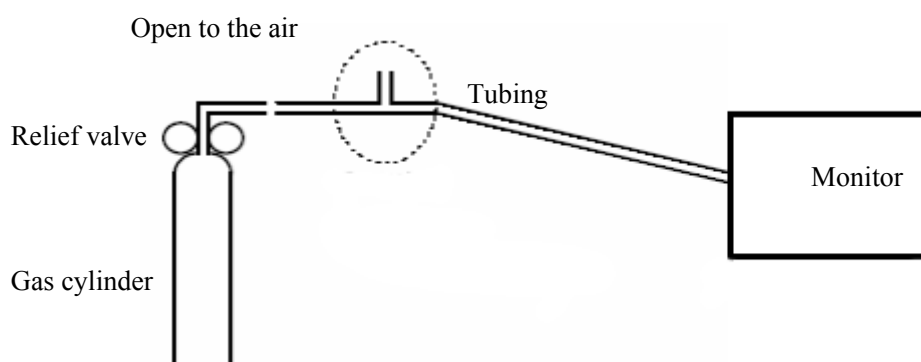
Calibration

Tools required:

- Gas cylinder, with a certain standard gas or mixture gas. Gas concentration should meet the following requirements: $AA \geq 1.5\%$, $CO_2 \geq 1.5\%$, $N_2O \geq 40\%$, $O_2 \geq 40\%$, of which AA represents an anesthetic agent. $a/c \leq 0.01$ (a is the gas absolute concentration accuracy; c is the gas concentration). For 100% O_2 calibration, a gas cylinder with 100% O_2 is used and the O_2 concentration is not less than 99%.
- T-shape connector
- Tubing

Follow this procedure to perform a calibration:

1. Select [**Main Menu**]→ [**Maintenance >>**]→ [**User Maintenance >>**]→ enter the required password→ [**Calibrate AG >>**].
2. Check the airway and make sure that there are no occlusions or leaks.
 - ◆ Vent the sampling tubing to the air and check if the [**Current FlowRate**] and [**Set FlowRate**] are approximately the same. If the deviation is great, it indicates that there is an occlusion in the tubing. Check the tubing for an occlusion.
 - ◆ Perform a leakage test to make sure that the airway has no leakage.
3. Connect the test system as follows:



4. Open the relief valve and vent a certain standard gas or gas mixture and make sure that there is an excess gas flow through the T-shape connector to air.
5. In the [**Calibrate AG**] menu, the concentration and flowrate of each measured gas are displayed.
 - ◆ If the difference between the measured gas concentration and the actual one is tolerable, a calibration is not needed.
 - ◆ If the difference is great, a calibration should be performed. Select [**Calibrate >>**] to enter the calibrate menu.

Calibrate AG		
CO2	0.0	%
N2O	0.0	%
O2	21.0	%
AA	---	%
Set FlowRate	120	ml/min
Current FlowRate	120	ml/min
<div>Calibrate >></div>		

6. Enter the vented gas concentration. If you use only one gas for calibration, set other gases' concentration to 0.
7. Select [**Start**] to start a calibration.
8. If the calibration is finished successfully, the message [**Calibration Completed!**] is displayed. If the calibration failed, the message [**Calibration Failed!**] is displayed. Perform another calibration.

CAUTION

- **Calibrate the O₂ module, If it has been transported for long distance or if you suspect it does not work properly.**
-

3.2.5 Preventative maintenance test report

Customer name			
Customer address			
Servicing person			
Servicing company			
Equipment under test (EUT)			
Model of EUT			
SN of EUT			
Hardware version			
Software version			
Test equipment	Model/No.	Effective date of calibration	
Test items	Test records	Test results (Pass/Fail)	
Visual inspection			

The case, display screen, buttons, knob, SMR, modules, power cord, wall-mount bracket and accessories have no obvious signs of damage.		
The external connecting cables are not frayed and the connector pins are not loose and bent.		
The external connectors are not loose or their pins are not bent.		
The safety labels and data plate are clearly legible.		
NIBP test		
The difference is within ± 3 mm when 0, 50 or 200 mmHg is set for NIBP accuracy test.		
There is no leakage with NIBP, or the manual leakage test result does not exceed 6mmHg/min.		
Sidestream CO₂ test		
Block the gas inlet of the module or watertrap. The sidestream CO ₂ flowrate is slower than 10ml/min and an alarm of CO ₂ Filterline Err is given. It indicates that there is no leakage.		
The displayed CO ₂ value is within $6 \pm 0.05\%$.		
Microstream CO₂ test		
Block the gas inlet of the module or watertrap. An alarm of CO ₂ Filterline Err is given. It indicates that there is no leakage.		
The displayed CO ₂ value is within $6 \pm 0.05\%$.		
AG test		
When AG flowrate is slower than 10ml/min, an alarm of AG Airway Occluded is given. It indicates that there is no leakage.		
The fan inside the AG module works properly.		
The measurement accuracy of CO ₂ , N ₂ O, O ₂ and AA (AA represents an anaesthetic agent) meets the product specifications in the Operator's Manual.		

3.3 Power On Test

This test is to verify that the patient monitor can power up correctly. The test is passed if the patient monitor starts up by following this procedure:

1. Insert two batteries in the battery chamber and connect the patient monitor to the AC mains, the AC mains LED and battery LED light.
2. Press the power on/off switch to switch on the patient monitor. The operating status LED lights up, and the technical and physiological alarm lamps light blue and red respectively.
3. After the start-up screens are displayed, the system sounds a beep indicating the self test on alarm sounds is passed. At the same time, the alarm lamp turns from yellow to red, and then turns off together with the technical alarm lamp. This indicates that the self test on alarm lamps is passed.
4. The patient monitor enters the main screen and start-up is finished.

3.4 Module Performance Tests

3.4.1 ECG Tests

ECG Performance Test

Tool required:

- Fluke Medsim 300B patient simulator recommended

Follow this procedure to perform the test:

1. Connect the patient simulator with the ECG module using an ECG cable.
2. Set the patient simulator as follows: ECG sinus rhythm, HR=80 bpm with the amplitude as 1mV.
3. Check the ECG waves are displayed correctly without noise and the displayed HR value is within 80 ± 1 bpm.
4. Disconnect each of the leads in turn and observe the corresponding lead off message displayed on the screen.
5. Set that the simulator outputs paced signals and set [**Paced**] to [**Yes**] on the monitor. Check the pace pulse marks on the monitor screen.

ECG Calibration

Tool required:

- Vernier caliper

Follow this procedure to perform a calibration:

1. Select the ECG parameter window or waveform area→ [**Filter**]→ [**Diagnostic**].
2. Select [**Main Menu**]→ [**Maintenance>>**].
3. Select [**Calibrate ECG**]. A square wave appears on the screen and the message [**ECG Calibrating**] is displayed.
4. Compare the amplitude of the square wave with the wave scale. The difference should be within 5%.
5. After completing the calibration, select [**Stop Calibrating ECG**].

If necessary, you can print out the square wave and wave scale through the recorder and then measure the difference.

3.4.2 Resp Performance Test

Tool required:

- Fluke Medsim 300B patient simulator recommended

Follow this procedure to perform the test:

1. Connect the patient simulator to the module using a non ESU-proof cable and set lead II as the respiration lead.
2. Configure the simulator as follows: lead II as the respiration lead, base impedance line as 1500 Ω ; delta impedance as 0.5 Ω , respiration rate as 40 rpm.
3. Check the Resp wave is displayed without any distortion and the displayed Resp value is within 40 ± 2 rpm.

3.4.3 SpO₂ Test

Tool Required:

- None.

Follow this procedure to perform the test:

1. Connect SpO₂ sensor to the SpO₂ connector of the monitor. Set [**Patient Cat.**] to [**Adu**] and [**PR Source**] to SpO₂ on the monitor.
2. Measure SpO₂ on your finger. (Assume that you stay healthy)
3. Check the Pleth wave and PR reading on the screen and make sure that the displayed SpO₂ is within 95%-100%.
4. Remove the SpO₂ sensor from your finger and make sure that an alarm of SpO₂ Sensor Off is triggered.

NOTE

-
- **A functional tester cannot be used to assess the accuracy of a pulse oximeter monitor. However, it can be used to demonstrate that a particular pulse oximeter monitor reproduces a calibration curve that has been independently demonstrated to fulfill a particular accuracy specification.**
-

3.4.4 NIBP Tests

See section 3.2.2 *NIBP Tests*.

3.4.5 Temp Test

Tool required:

- Resistance box (with accuracy above 0.1Ω)

Follow this procedure to perform the test:

1. Connect the two pins of any Temp connector of a module to the two ends of the resistance box using 2 wires.
2. Set the resistance box to 1354.9Ω (corresponding temperature is 37°C).
3. Verify each Temp channel of the monitor and make sure that the displayed value is within $37 \pm 0.1^{\circ}\text{C}$.

You can also use a patient simulator to perform the Temp test.

3.4.6 IBP Tests

IBP Performance Test

Tool required:

- Medsim300B patient simulator, MPS450, or other equivalent device
- Dedicated IBP adapter cable (300B, P/N 00-002199-00) (use P/N 00-002198-00, if the simulator is MPS450)

Follow this procedure to perform the test:

1. Connect the patient simulator with the pressure module.
2. Make the patient simulator outputs 0 to each IBP channel.
3. Press the Zero Key on the module to make a zero calibration.
4. Configure the patient simulator as P (static) = 200 mmHg.
5. The displayed value should be within 200 ± 2 mmHg.
6. If the value is outside of these tolerances, calibrate the pressure module. If the IBP module was calibrated with a dedicated reusable IBP sensor, check the calibration together with this IBP sensor.
7. Make that the patient simulator outputs 120/80 mmHg ART signals and 120/0 mmHg LV signals respectively to each IBP channel and check that the IBP wave is displayed correctly.
8. Repeat the steps above for all the IBP channels.

IBP Pressure Calibration

Method 1:

Tools required:

- Medsim300B Patient simulator, MPS450, or other equivalent device
- Dedicated IBP adapter cable (300B, P/N 00-002199-00) (use P/N 00-002198-00, if the simulator is MPS450)

Follow this procedure to perform the test:

1. Connect the patient simulator to the pressure connector on the module.
2. Set the patient simulator to 0 pressure for the desired IBP channel.
3. Press the Zero Key on the module to make a zero calibration.
4. Configure the patient simulator as P (static) = 200 mmHg.
5. Select [Main Menu] → [Maintenance >>] → [User Maintenance >>] → [Cal. IBP Press. >>]. In the [Cal. IBP Press.] menu, set the calibration value to 200 mmHg.

-
6. Select the [**Calibrate**] button next to the desired IBP channel to start a calibration.
 7. If the calibration is completed successfully, the message [**Calibration Completed!**] will be displayed. Otherwise, a corresponding message will be displayed.

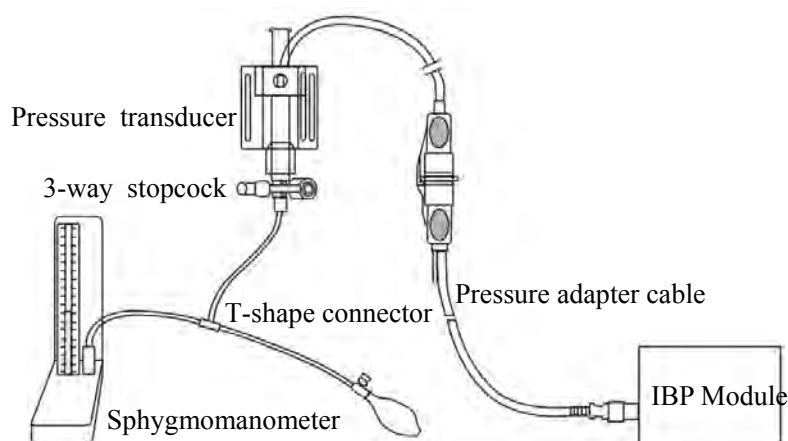
Method 2:

Tools required:

- Standard sphygmomanometer
- Balloon pump
- Tubing
- T-shape connector

To perform a calibration:

1. Connect the 3-way stopcock, the sphygmomanometer and the balloon pump through a T-shape connector, as shown below.
2. Zero the transducer. Then open the stopcock to the sphygmomanometer.
3. Press the Main menu button on the equipment's front panel. Select [**Maintenance>>**]→ [**User Maintenance >>**]→ enter the required password→ [**Cal. IBP Press. >>**]. Then configure IBP calibration value.
4. Inflate using the balloon pump until the reading of sphygmomanometer approximates the preset calibration value.



5. Adjust the calibration value in the [**Cal. IBP Press.**] menu until it is equal to the reading of sphygmomanometer
6. Select the [**Calibrate**] button to start a calibration
7. The message [**Calibration Completed!**] is displayed after a successful calibration. If the calibration failed, the prompt [**Calibration Failed!**] will be displayed.

3.4.7 C.O. Test

Tools required:

- Medsim300B Patient simulator
- C.O. adapter box

Follow this procedure to perform the test:

1. Connect the patient simulator to the C.O. module using a C.O. main cable.
2. Set the blood temperature (BT) to 37°C on the patient simulator and check the temperature value is $37 \pm 0.1^\circ\text{C}$.
3. Set [**Auto IT**] to [**Off**] and adjust [**IT**] to 24°C. Select [**C.O. Measure**] to enter the C.O. measurement window and set [**Comp. Const.**] to 0.595.
4. Set the injectate temperature to 24°C and the C.O. to 5L/min on the C.O. simulator. Select [**Start**] in the C.O. measurement window to start C.O. measurements and after 3-10 seconds press the run key on the simulator.
5. Check the C.O. value is $5 \pm 0.25\text{L/min}$.

3.4.8 Mainstream CO₂ Tests

NOTE

-
- **Make sure that the barometric pressure set in [Maintain CO₂>>] of [User Maintenance>>] accords with the local barometric pressure before performing mainstream CO₂ tests.**
-

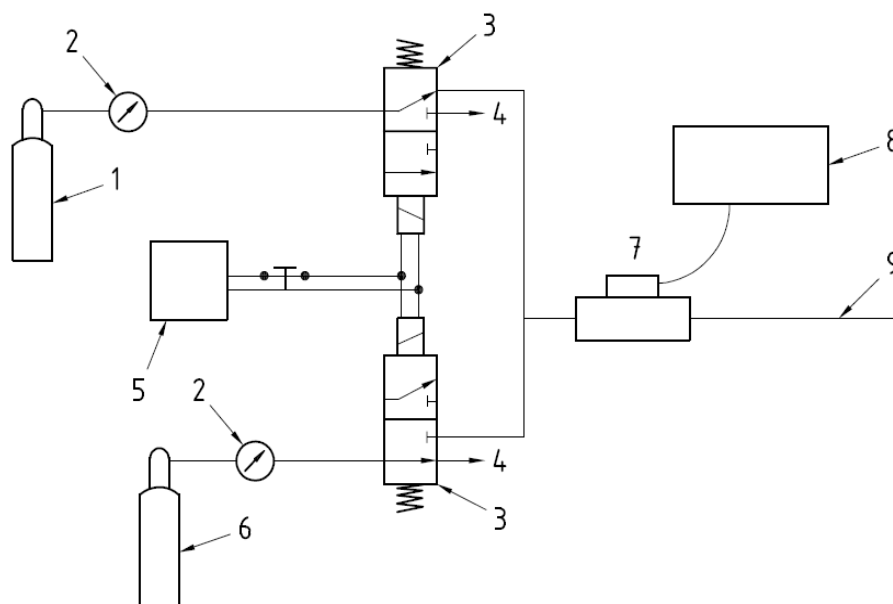
Tools required:

- A steel gas cylinder with $6 \pm 0.05\%$ CO₂
- A steel gas cylinder with compressed air or N₂ (with standard concentration)
- Two 3-way valves (power supply controlled)
- Flowmeter
- Power supply
- Tube

Follow this procedure to perform the test:

1. Wait until CO₂ warmup is finished and then select [**Start Zero Cal.**] from [**CO₂ Setup**] menu to start a zero calibration. If the zero calibration fails, the prompt message [**CO₂ Zero Failed**] is displayed. Otherwise, the baseline of waveform recovers to zero.
2. Set [**Apnea Time**] to 10 s in the [**Adjust CO₂ Limits**] menu.

- 3 Blow to the CO₂ sensor to generate a CO₂ waveform and then place the sensor in the air. Check if the alarm message [CO₂ Apnea] is displayed on the screen.
- 4 Connect the test system as follows



Indication of numbers in the figure above

1	A steel gas cylinder with $6 \pm 0.05\%$ CO ₂
2	Flowmeter
3	3-way valve (power supply controlled)
4	Open to air
5	Power supply (controlling two 3-way valves)
6	Compressed air or N ₂ with standard concentration
7	Mainstream CO ₂ sensor
8	Patient monitor
9	Tube (preventing back flow)

- 5 Adjust the power supply and turn on/off 3-way valves to ensure that that only one cylinder is connected to Mainstream CO₂ sensor via 3-way valves at one time and the flowmeter reading is stable and within 2-5L/min.
- 6 Switch between the two cylinders to connect Mainstream CO₂ sensor at intervals of 6 -10s and check if the displayed CO₂ value is within $6 \pm 0.05\%$.

3.4.9 Sidestream and Microstream CO₂ Module Tests

See section 3.2.3 *Sidestream and Microstream CO₂ Module Tests*.

3.4.10 AG Tests

See section 3.2.4 *AG Tests*.

3.4.11 ICG Test

Tool required:

- ICG simulator (BZ-4575)

1. Connect the ICG simulator and the patient monitor using standard ICG cable and sensor.
2. Select **[ICG setup]**→ **[Patient Demographics >>]** and then input parameter values as follows:

Height	180 cm	CVP	6 mmHg
Weight	75 kg	PAmean	8 mmHg
PAWP	10 mmHg		

3. Switch on the simulator and set as follows: HR=60±1 bpm, VI=61±4/1000s, TFC=32±2/kOhms. Then, start ICG monitoring.
4. After the measurement becomes stable and check that the measured results are as follows: HR=60±2 bpm, VI=61±4/1000s, TFC=32±2/kOhms.
5. Set on the simulator as follows: HR=70±1 bpm, VI=48±4 /1000s, TFC=32±2 / kOhms, and then start ICG monitoring. After the measurement becomes stable and check that the measured results are as follows: HR=70±2 bpm, VI=48±4 /1000s, TFC=32±2 / kOhms.

3.4.12 BIS Test

You can choose either of the following methods to perform the test:

Method 1:

Tools required:

- None.

1. Connect the BIS sensor to a healthy, wide-awake adult as directed in the Operator's Manual.
2. Check the EEG wave and BIS numerics displayed on the screen and make sure the BIS value is within 80-100.

Method 2:

Tools required:

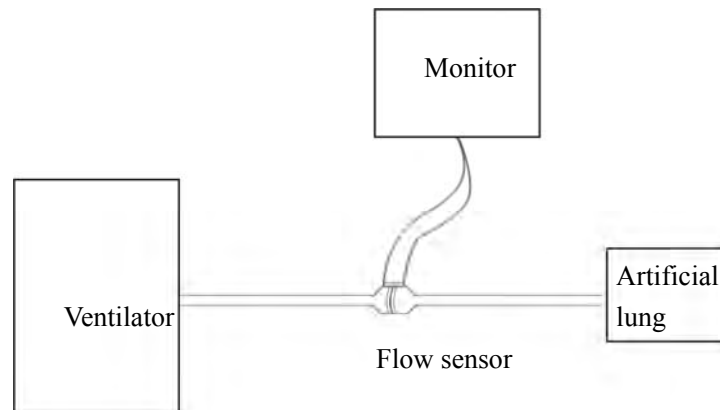
- BIS simulator (with flexible cable)

1. Connect the BIS sensor with the BIS simulator and select [**BIS Setup**]→ [**BIS Sensor Check**] to perform a cyclic impedance check.
2. After the cyclic impedance check is finished, check that the result for each electrode is pass.
3. Check the EEG wave and BIS numeric displayed on the screen.

3.4.13 RM Test

Tool required:

- Gas source
- Ventilator (calibrated)
- Artificial lung
- Pediatric/neonate flow sensor



Follow this procedure to perform the test:

1. Connect the equipment as shown above. Make sure that the blue sensing tube on the flow sensor is connected with the artificial lung.
2. Set [**Patient Cat.**] to [**Adu**]. In the [**RM Setup**] menu, select [**Sensor Type**] according to the used sensor and set [**Ventilation Mode**] to [**Mechanical**].
3. Enter the [**RM Setup**] menu and select [**Calibrate >>**]. Input the constant marked on the sensor and calibrate the flow sensor.
4. Configure the ventilator as follows: TV=500 ml, RR =20 rpm, I:E=1:2.
5. Select [**Respiratory Loop**] in the [**RM Setup**] menu. Verify that the displayed TV is within 500±50ml and RR is within 20±1rpm.

3.4.14 CCO/SvO₂ Tests

Interconnecting Function

Tools required:

- None.

1. Connect and set the patient monitor and Vigilance monitor per the procedures in the Operator's Manual.
2. Set the Vigilance monitor to Demo mode.
3. Check that the CCO/SvO₂ numerics displayed on the patient monitor and Vigilance monitor are consistent.

Output Performance

Tools required:

- Oscilloscope

1. Connect the signal output end of the connecting cables of the CCO/SvO₂ module to the oscilloscope.
2. Make the monitor to perform an ECG calibration. Check that the ECG waves displayed on the oscilloscope are consistent with the ECG calibration waves displayed on the monitor screen.
3. Select [**CCO Setup**]→ [**Signal Output Setup >>**] and then select [**Simulated High Value**] from the pop-up menu. Check that the amplitude of electrical level at the signal output port of MAP, CVP and SpO₂ are $5\pm0.25V$, $5\pm0.25V$ and $10\pm0.5V$ respectively.

3.4.15 PiCCO Tests

Performance Test

Tool required:

- Medsim300B patient simulator
- PiCCO IBP test cable (6800-J87)

Follow this procedure to perform the test:

1. Connect the patient simulator, the PiCCO IBP test cable and the PiCCO module.
2. Let the patient simulator outputs 0 mmHg respectively to the pArt channel and the pCVP channel.
3. In the **[pArt Setup]** menu, select **[pArt Zero >>]→[Zero]**.
4. In the **[pCVP Setup]** menu, select **[pCVP Zero >>]→[Zero]**.
5. Let the patient simulator output static pressure 200 mmHg to pArt channel and 20 mmHg to pCVP channel.
6. The pArt value displayed on the monitor should be within 200 ± 4 mmHg, and pCVP value within 20 ± 1 mmHg.
7. If the pArt error is beyond ± 4 mmHg or pCVP error beyond ± 1 mmHg, calibrate the PiCCO module. If the module was calibrated with a dedicated reusable IBP sensor, check the calibration together with this IBP sensor.
8. Let the patient simulator output ART signal to the pArt channel and pCVP signal to the pCVP channel, verify that the pArt and pCVP waves are displayed correctly.

Pressure Calibration

Method 1

Tools required:

- Medsim300B patient simulator
- PiCCO IBP test cable (6800-J87)

Follow this procedure to perform the test:

1. Connect the patient simulator, the PiCCO IBP test cable and the PiCCO module.
2. Let the patient simulator outputs 0 mmHg respectively to the pArt channel and the pCVP channel.
3. In the **[pArt Setup]** menu, select **[pArt Zero >>]→[Zero]**.
4. In the **[pCVP Setup]** menu, select **[pCVP Zero >>]→[Zero]**.
5. Set static pressure to 200 mmHg on the patient simulator.

-
6. Select [**Main Menu**]→[**Maintenance >>**]→[**User Maintenance >>**]→ [**Cal. IBP Press. >>**]. In the [**Cal. IBP Press.**] menu, set the calibration pressure to 200 mmHg.
 7. Select the [**Calibrate**] button next to the desired IBP channel to start a calibration.

If the calibration is completed successfully, the message [**Calibration Completed!**] will be displayed. Otherwise, a corresponding message will be displayed.

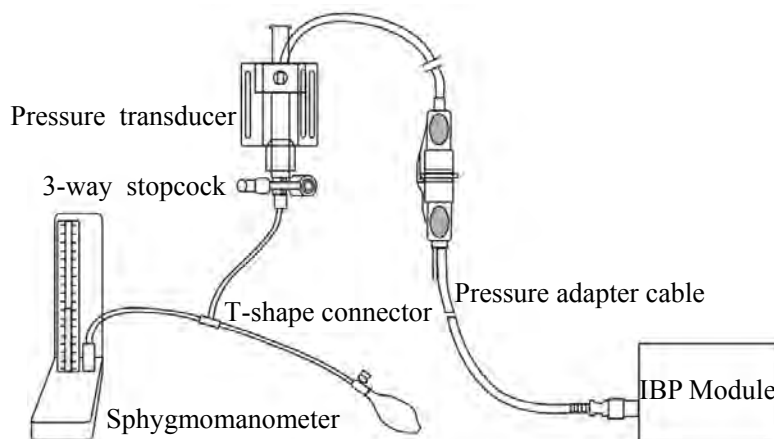
Method 2

Tools required:

- Standard sphygmomanometer
- Balloon pump
- Tubing
- T-shape connector

To perform a calibration:

1. Connect the 3-way stopcock, the sphygmomanometer and the balloon pump through a T-shape connector, as shown below.
2. Vent the transducer to the atmospheric pressure by turning on the 3-way stopcock to the air. Zero the transducer, and then open the stopcock to the sphygmomanometer.
3. Select [**Main Menu**]→[**Maintenance >>**]→ [**User Maintenance >>**]→enter the required password→ [**Cal. IBP Press. >>**]. In the [**Cal. IBP Press.**] menu, set the calibration pressure to 200 mmHg.
4. Inflate using the balloon pump until the reading of sphygmomanometer approximates the preset calibration value.



5. Adjust the calibration value in the [**Maintain IBP**] menu until it is equal to the reading of sphygmomanometer
6. In the [**Cal. IBP Press.**] menu, select the [**Calibrate**] button next to the desired IBP channel to start a calibration

The message [**Calibration Completed!**] is displayed after a successful calibration. If the calibration failed, the prompt [**Calibration Failed!**] will be displayed.

C.O. Test

Tools required:

- Medsim300B Patient simulator, or equivalent equipment
- C.O. adapter box (for 300B)
- PiCCO test cable (PN: 040-001301-00)

Follow this procedure to perform the test:

1. Connect the patient simulator and the C.O. module using a C.O. trunk cable and a C.O. adapter box.
2. Set the blood temperature (BT) to 37°C on the patient simulator and check the temperature value displayed on the monitor is $37 \pm 0.1^\circ\text{C}$.
3. In the [**CCO Setup**] menu, select [**PiCCO Guide>>**], and check if the value for [**Cat.Type**] is [PV2015L20].
4. Turn the injectate temperature (TI) knob on the C.O. adapter box to set the TI to $20 \pm 1^\circ\text{C}$ for the patient simulator and C.O. to 5L/min.
5. In the PiCCO screen, select [**Start**] to start C.O. measurement. As soon as the prompt [**Inject XXml**] is displayed, adjust TI to $4 \pm 1^\circ\text{C}$, and then quickly back to $20 \pm 1^\circ\text{C}$. Simultaneously press the button on the simulator that corresponds to 5L/min. The whole procedure shall be finished in 10 s.
6. Verify that the C.O. value displayed on the monitor is correct.

3.4.16 ScvO₂ Tests

You can perform ScvO₂ test by either of the following two methods:

Method 1:

Tools required:

- PC (installed OM_Simulation software, G-6800-S15)
- ScvO₂ test cable (6800-J68)

1. Connect the ScvO₂ test cable to the ScvO₂ module and the serial port on the PC respectively;
2. Open ScvO₂ test software to select the corresponding serial port from [**COM port**] menu (default serial port: COM1);
3. Select [**Start**] from the [**Replay**] menu. Then the ScvO₂ measurement area is displayed on the patient monitor;
4. Enter [**ScvO₂ Setup**] menu and set [Hb/Hct] to [Hct];
5. Set [**LED660**], [**LED800**], [**LED880**] to '1250, 40000' (inching with the direction key on the keyboard if needed). Select [**Start**] from the [**Replay**] menu;
6. Enter [**ScvO₂ Calibration**] menu and select [**Sample drawn**] when SQI occupies four grids (80%). In the popup menu, set [**ScvO₂**] and [**Hct**] to 67% and 40% respectively, then select [**Calibrate**]. Close [**ScvO₂ Calibration**] menu to check that the ScvO₂ value is displayed as 67% in the monitoring screen.
7. Set the value of [**LED660**] to '1250, 33600' and keep other LCD values unchanged. Check that the ScvO₂ measurement is within $(50 \pm 3)\%$.
8. Set the value of [**LED660**] to '1250, 46400' and keep other LCD values unchanged. Check that the ScvO₂ measurement is within $(80 \pm 3)\%$.

Method 2:

Tools required:

- None.

1. Connect the ScvO₂ sensor to the patient monitor. Check that the front end of the ScvO₂ sensor illuminates normally.
2. Nip the front end of the ScvO₂ sensor with two fingers to perform a ScvO₂ calibration.
3. Check that the patient monitor displays the ScvO₂ measurement normally.

3.4.17 NMT Tests

NMT Performance Test

Tool required:

- Resistance box
- Oscilloscope (Agilent DSO6052A)

1. Set up the resistance box:
 - a. Set the resistance value to 1kOhm.
 - b. Connect the stimulation electrodes to the two wiring terminals.
2. Connect the oscilloscope sensors to the NMT stimulation electrodes, making sure that each sensor and electrode connected have the same polarity.
3. Set up the monitor:
 - a. Insert the NMT module into the module rack of the monitor.
 - b. Set the [**Stimulation Current**] to [**Supra(60mA)**].
 - c. Set the [**Pulse Width**] to 300μs.
 - d. Perform an ST measurement.
4. Proceed as follows to capture the pulse signals and measure the [**PK-PK**] and [**+Width**]. The following procedures take the oscilloscope of Agilent DSO6052A as an example.
 - a. Power on the oscilloscope by pressing the [**POWER**] button. Select the button [**1**] and verify the button light is on.
 - b. Press the [**Mode/Coupling**] button, and select [**Normal**], [**Noise REJ**] and [**HF Reject**] at the lower part of the screen.
 - c. Press the [**Acquire**] button, and select [**High Resolution**] from the dropdown list of [**Acq Mode**].
 - d. Press the [**Quick Meas**] button. Select [**PK-PK**] from the dropdown list of [**Select**] and select [**Measure**] to confirm; select [**+Width**] from the dropdown list of [**Select**] and then select [**Measure**] to confirm.
 - e. In the [**Horizontal**] area, adjust [**Delay**] to 0.0s by using the [**◀▶**] knob, and adjust the numeric value to 100ms by using the big knob.
 - f. In the [**Analog**] area, adjust the numeric value to 20.0V by using the big knob and adjust the [**Ch(1)**] to 40.0V by using the [**◆**] knob.
 - g. Press the [**Edge**] button. Select [**1**] from the dropdown list of [**Source**]; select [**Rising**] from the dropdown list of [**Slope**].
 - h. In the [**Trigger**] area, adjust the numeric value to 40.0V by using the [**Level**] knob. Press [**Single**] to perform a measurement.



The test passes when the measurements are within the following ranges:

Test items	Measurements
【PK-PK】	54V - 66V
【+Width】	270 μ s - 330 μ s

Checking NMT Sensor

Tools required:

- None

1. Connect the patient monitor, NMT module, and NMT accessories.
2. Select [Main Menu]→ [Maintenance >>]→ [User Maintenance >>]→ enter the required password→ [NMT Sensor Check >>] from the patient monitor.
3. Follow the on-screen instructions to check the NMT sensor in four ways.

If sensor check completes successfully, the message “Test passed. The function of NMT sensor is OK” is presented. If any of the four steps fails, check if the sensor is placed correctly as instructed, and perform the sensor check again. Replace the sensor if you cannot pass the sensor check..

NOTE

- Stop NMT measurement or calibration before starting NMT sensor check.
- Take care to handle the the NMT sensor. Avoid forcefully striking the sensor.

3.4.18 EEG tests

Method 1

Tools required:

- EEG simulator (Nannini 169/1)

1. Insert the EEG module into the module slot of the monitor.
2. Connect the EEG module to the EEG simulator through the EEG patient cable.
3. Select **[EEG Setup]**→ **[Sensor Check]** to perform a sensor check.
4. Check that the result for each electrode is pass.
5. Check that the EEG wave and numeric are displayed on the screen.

Method 2

Tools required:

- None

1. Insert the EEG module into the module slot of the monitor.
2. Connect the EEG cable and EEG electrodes to the EEG module, and short all the electrodes.
3. Select **[EEG Setup]**→ **[Sensor Check]** to perform a sensor check.
4. Check that the result for each electrode is pass.
5. Check that the EEG wave and numeric are displayed on the screen.

3.5 Nurse Call Relay Performance Test

Tools required:

- Multimeter

1. Connect the nurse call cable to the Nurse Call Connector of the patient monitor.
2. Enter Demo mode. Then, select [**Main Menu**] → [**Maintenance >>**] → [**User Maintenance >>**] → enter the required password → [**Others >>**] → [**Auxiliary Output**] → [**Nurse Call**].
3. In the [**Others >>**] menu, select [**Nurse Call Setup >>**] and then select all options of [**Alm Lev**] and [**Alarm Cat.**] and set [**Contact Type**] to [**Normally Open**]
4. In [**Nurse Call Setup >>**] setup menu, set [**Signal Type**] to [**Pulse**]. Make the monitor to generate an alarm and check that the output are pulses of 1s width and the relay contacts are close (can be measured with a multimeter) when there is an alarm.
5. In [**Nurse Call Setup >>**] setup menu, set [**Signal Type**] to [**Continuous**]. Make the monitor to generate an alarm and check that the output is continuous high level and the relay contacts are close (can be measured with a multimeter) when there is an alarm.

3.6 Analog Output Performance Test

Tool required:

- Patient simulator
- Oscillograph

1. Connect the patient simulator to the monitor using an ECG or IBP cable and connect the oscillograph to the Auxiliary Output Connector of the patient monitor.
2. Select [**Main Menu**] → [**Analog Output Setup**]. Switch Analog Output [**On**].
3. Verify that the waves displayed on the oscillograph are identical with those displayed on the monitor.



3.7 Electrical Safety Test

See *A Electrical Safety Inspection* for electrical safety tests.

3.8 Touchscreen Calibration

Tools required:

- None.

1. Select the [**Cal. Screen**] QuickKey or select [**Main Menu**]→ [**Maintenance >>**]→ [**User Maintenance >>**]→ enter the required password→ [**Cal. Touchscreen**].
2. The  symbol will appear at different positions of the screen.
3. Select, in turn, the central point of the  symbol.
4. After the calibration is completed, the message [**Screen Calibration Completed!**] is displayed. Select [**Ok**] to confirm the completion of the calibration.

3.9 Recorder Check

Tools required:

- None.

1. Print ECG waveforms. The recorder should print correctly and the printout should be clear.
2. Set the recorder to some problems such as out of paper, etc. the patient monitor should give corresponding prompt messages. After the problem is removed, the recorder should be able to work correctly.
3. Switch automatic alarm recording for each parameter ON and then set each parameter's limit outside set alarm limits. Corresponding alarm recordings should be triggered when parameter alarms occur.

3.10 Network Print Test

Note

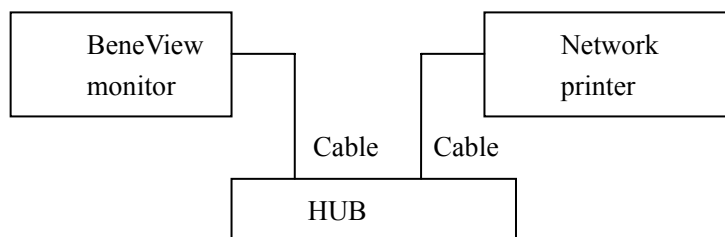
-
- **HP LaserJet 1505n, P2035n, P4015n, or 1606dn laser printer is recommended for BeneView series of patient monitors.**
-

Tools required:

- Hub and network cable

3.10.1 Equipment Connection and Setup

1. Connect the patient monitor and network printer to a HUB using common network cables as follows:



2. Set IP address as follows: Select [**Main Menu**]→ [**Maintenance >>**]→ [**User Maintenance >>**]→ enter the required password→ [**IP Address Setup >>**] and set the IP address of the patient monitor in the same network segment with that of the network printer. (See the instructions for use accompanying the printer)
3. Search for printer by selecting [**Main Menu**]→ [**Print Setup >>**]→ [**Printer Setup >>**]→ [**Search Printer**]. After a while, the printer's model and IP address will appear in the box beside [**Printer**].

3.10.2 Print Function Test

1. Enter the Demo mode of the patient monitor.
2. Select [**Main Menu**]→ [**Print Setup >>**]→ [**Realtime Reports >>**]→ [**Normal Report**] and then select [**Print**]. The network printer shall print out the report correctly.

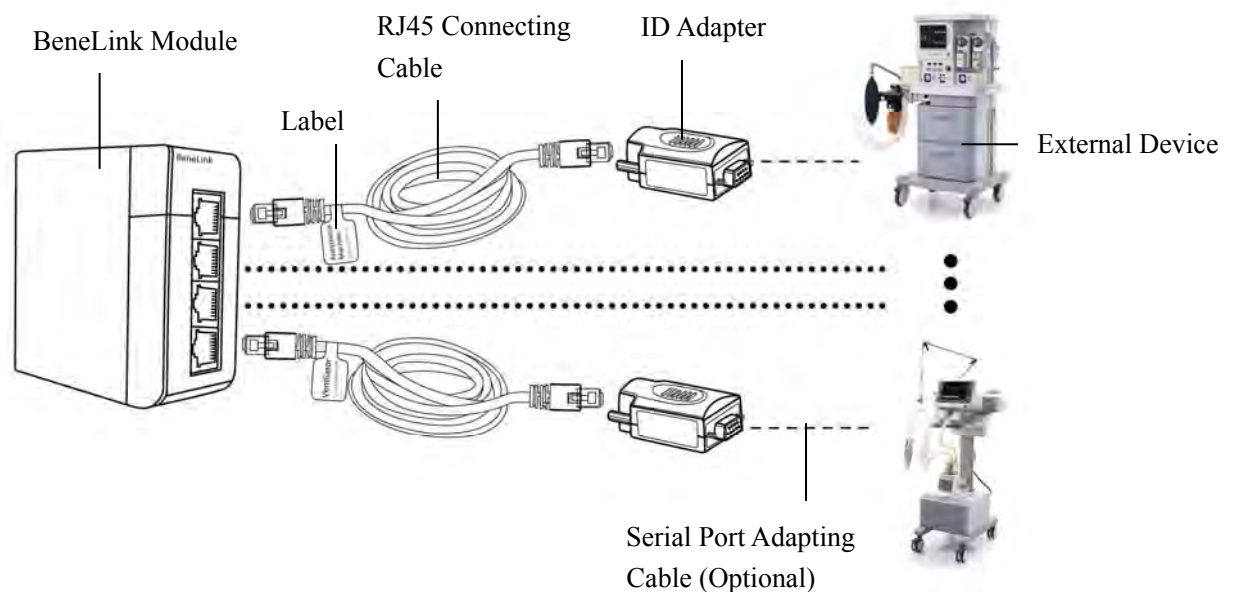
3.11 BeneLink Module Check

3.11.1 Device Connection and Setup

Tools required:

- External device (anesthesia machine, ventilator)
- ID adapter that matches the external device
- RJ45 connecting cable
- Serial port adapting cable that matching the external device

Please refer to the following procedure to connect an external device:



1. Insert the BeneLink module into the module slot on the BeneView patient monitor.
2. Connect the ID adapter that matches the external device to the BeneLink module with a RJ45 connecting cable.
3. Plug the ID adapter into the RS232 port on the external device. Some external devices may have ports incompatible with the ID adapter. In this case, a serial port adapting cable is required. Please be sure that you have selected the proper cable before connection.
4. Stick a label indicating device name to the RJ45 connecting cable at the end nearby the BeneLink module. When the BeneLink module is connected to several external devices, you can tell the devices apart easily with these labels.
5. Switch the external device on.

NOTE

- **Devices of the same category can not be connected to the BeneLink module simultaneously.**
 - **Use the serial port adapting cable only with its matching external device. Please see the following table to select the correct adapting cable.**
 - **Use the ID adapter only with the matching external device. Please see the following table for correct ID setup in [Factory Maintenance] menu.**
-

External Device	ID for ID adapter	Type of Serial Port Adapting Cable
Mindray Wato 20/30/55/65	4D52B2AE	No need to use the adapting cable: the ID adapter can be plugged into the serial port of the external device directly.
Newport E360	4E50B1B0	Type B
Puritan Bennett 840	SNDF: 5042AFBE (recommended) SNDA: 5031AFCF (support less parameters than protocol SNDF)	No need to use the adapting cable: the ID adapter can be plugged into the serial port of the external device directly.
Maquet Flow-i	4D46B2BA	Type B
Maquet Servo-i/Servo-s	4D53B2AD	Type B
Draeger Evita 2 / Evita 2 dura / Evita 4/ Evita XL	4434BBCC	Type B
Hamilton G5 (protocol Polling)	3550CAB0	Type B
Hamilton C2 (protocol Polling)	3270CD90	Type B
Hamilton Galileo (protocol Polling)	4750B8B0	Type B
Hamilton G5 (protocol Block)	3542CABE	Type B
Ohmeda Avance/Aisys	4F41B0BF	Type D
Ohmeda Aestiva 7100/7900	4F37B0C9	Type D

External Device	ID for ID adapter	Type of Serial Port Adapting Cable
Drager Fabius GS/Plus/Trio	4446BBBA	GS: no need to use the adapting cable: the ID adapter can be plugged into the serial port of the external device directly. Plus: type C Trio: type C
Drager Primus	4450BBB0	Type C
TCM CombiM/TCM TOSCA	5443ABBD	Type C
TOF-Watch SX	5457ABA9	Type C
Expand Model	/	Type A

3.11.2 Device Integration Function Test

3.11.2.1 Preparation

Prepare the tools needed for function test according to the type of the external device you install. Please see the Instructions for Use of the corresponding external device for guidance. For the function test of ventilator and anesthesia machine, at least the following tools are needed:

- BeneView patient monitor with BeneLink module properly installed
- External device (anesthesia machine or ventilator) under test
- Gas source(tube or gas cylinder), including air or O₂ at least, and N₂O or other anesthesia gases are optional
- Tube that connects the patient(or test lung)
- Test lung and a matching Y-pipe, or other accessories

3.11.2.2 Procedure and Items to Be Checked

Follow the steps below:

1. Connect the BeneLink module to the ventilator or the anesthesia machine. Please see Device Connection and Setup for more details.
2. Connect the gas supply and test lungs to the ventilator or anesthesia machine, turn on the device, and configure as follows:
 - ◆ Setup up the serial port of the external device by referring to *Serial Port Configuration List*.

-
- ◆ Setup up the pressure control mode and check if the ventilator or anesthesia machine works normally.
3. Check the ID adapter is correctly configured, and the green indicator of corresponding port on the BeneLink module illuminates constantly.
 4. Access [**Devices Integrated**] screen on the patient monitor. Check that the device type (ventilator or anesthesia machine) and ventilation mode are correctly displayed.
 5. Select parameters PEEP, Pmean, VTe, MV, I:E, and f(RR) respectively on the patient monitor and check if the parameter values displayed on the patient monitor are consistent with those displayed on the ventilator or anesthesia machine.
 6. Re-configure the above parameters on the ventilation or the anesthesia machine and check if the parameter values displayed on the patient monitor change accordingly.
 7. Trigger alarms [**MV Too Low**], [**Airway Pressure Too High**], [**PAW Too High**], [**Peak Too High**], and [**No Gas Supply**] (no Air or O2) on the ventilation or the anesthesia machine. Check that these alarm messages are correctly recorded in the alarm list of the patient monitor.
 8. Switch the ventilator or anesthesia machine to volume control ventilation mode. Check if the ventilation mode displayed on the patient monitor changes accordingly, and if the parameter values of PEEP, Pmean, VTe, MV, I:E, and f(RR) are correctly displayed.

Serial Port Configuration List

External Device	Setup	Remark
Mindray Wato 20/30/55/65	Not required.	If you need to view the parameters of CO2、AG、BIS module in the anesthesia machine, select [Factory Mainenance>>] → [Function Configuration>>] → [Select Module] in standby mode and tick the corresponding module.
Maquet Servo-i Maquet Servo-s	Not required.	The following information is for further reference: Baud Rate: 9600 bps Word Length: 8 bits Parity: even Stop Bits: 1
Draeger Evita 2	Channel A: Not required; Channel B: Protocol: Medibus Baud rate: 19200 Parity: even Stop Bits: 1	/

External Device	Setup	Remark
Draeger Evita 2 dura/ Evita 4/ Evita XL	Protocol: Medibus Baud Rate: 19200 Parity: even Stop Bits: 1 Interval: ---(Evita 2 dura)	/
Newport E360	Protocol: Newport	The following information is for further reference: Baud Rate: 38400 bps Word Length: 8 bits Parity: NONE Stop Bits: 1
Puritan Bennett 840	Baud Rate: 38400 Word Length: 8 bits Parity: NONE	/
Hamilton G5 (protocol Polling)	Protocol: Polling.	The following information is for further reference: Baud Rate: 9600 bps Word Length: 7 bits Parity: even Stop Bits: 2
Hamilton C2 (protocol Polling)	Protocol: Polling.	The following information is for further reference: Baud Rate: 9600 bps Word Length: 7 bits Parity: even Stop Bits: 2
Hamilton Galileo (protocol Polling)	Not required.	The following information is for further reference: Baud Rate: 9600 bps Word Length: 7 bits Parity: even Stop Bits: 2
Hamilton G5 (protocol Block)	Protocol: Block.	The following information is for further reference: Baud Rate: 38400 bps Word Length: 8 bits Parity: none Stop Bits: 1

External Device	Setup	Remark
Ohmeda Avance/Aisys	Not required.	The following information is for further reference: Baud Rate: 19200 bps Word Length: 7 bits Parity: odd Stop Bits: 1
Ohmeda Aestiva 7100 /7900	Not required.	The following information is for further reference: Baud Rate: 19200 bps Word Length: 7 bits Parity: odd Stop Bits: 1
Drager Fabius GS/Plus/Tiro	Protocol: Medibus Baud Rate: 9600 Word Length: 7 bits Parity: even Stop Bits: 1	/
Drager Primus	Protocol: Medibus Baud Rate::9600 Word Length: 8 bits Parity: even Stop Bits: 1	/
TCM CombiM/TCM TOSCA	Protocol: Monlink.	The following information is for further reference: Baud Rate: 9600 bps Word Length: 8 bits Parity: even Stop Bits: 1
TOF-Watch SX	Not required.	The following information is for further reference: Baud Rate: 19200 bps Word Length: 8 bits Parity: none Stop Bits: 1

3.11.3 Installation and Test Report

Basic Information			
Hospital		Department	
Serial number of ID adapter		Name of the external device	
ID of the external device		Type of serial port adapting cable	
Software version and other information of the external device			
Checking the connection			Test Result (Yes/No)
Can the patient monitor and the external device be assembled together using designated accessories?			
Does the green indicator of corresponding port on the BeneLink module illuminate while the other indicators are not?			
Are there numerics or characters displayed on the [Devices Integrated] screen of the patient monitor?			
Are the device type and parameter values displayed correctly on the [Devices Integrated] screen of the patient monitor when the external device just enters working mode?			
Is the ventilation mode correctly displayed on the patient monitor? Does it change correctly when the ventilation mode on the external device is changed?			
Checking the parameters			
Parameters on the external device	Value	Parameters on the patient monitor	Value
PEEP		PEEP	
Pmean		Pmean	
VT _e		VT _e	
f (RR)		f (RR)	
P _{peak}		P _{peak}	
Checking the alarms			
Alarm	Alarm messages displayed on the external device	Alarm messages displayed in the alarm list of the patient monitor	
RR Too High			
Apnea			
Patient Disconnected			

Other information

3.12 Battery Check

Tools required:

- None.

Function Test

1. If the patient monitor is installed with batteries, remove the batteries first.
2. Verify that the patient monitor works correctly when running powered form an AC source.
3. Insert two batteries per the procedures provided in the Operator's Manual.
4. Remove the AC power cord and verify that the patient monitor still works correctly.
5. For T5 only: Remove one battery and verify that the patient monitor continues to work correctly. Verify that the patient monitor can also work independently from another battery.

Performance Test

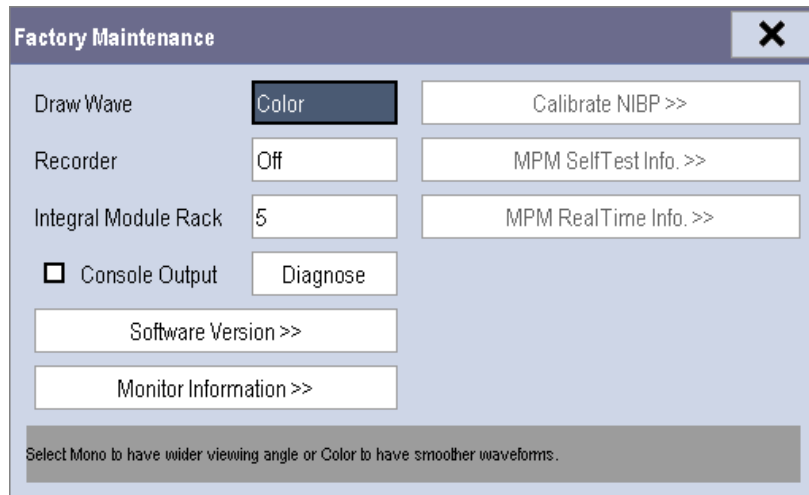
Perform the test by referring to the **Battery** chapter in the Operator's Manual and verify the operating time of the battery meets the product specification.

3.13 Factory Maintenance

3.13.1 Accessing Factory Maintenance Menu

To access the factory maintenance menu, select **[Main Menu]**→ **[Maintenance >>]** → **[Factory Maintenance]** and then enter the required password.

The **[Factory Maintenance]** menu is shown below.



3.13.2 Drawing Waves

There are two methods to draw waves: Color and Mono.

- Color: selecting Color will have smoother waveforms.
- Mono: selecting Mono will have a wider viewing angle.

3.13.3 Recorder

To enable/disable the recorder, select **[Recorder]** and toggle between **[On]** and **[Off]**.

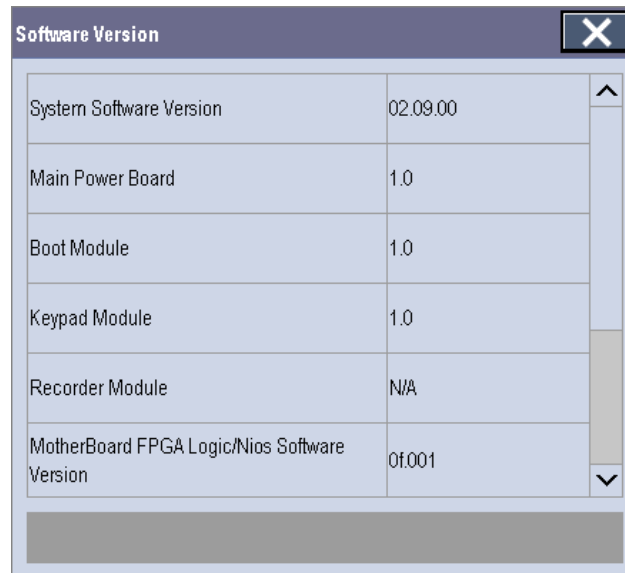


CAUTION

- The recorder is disabled if **[Recorder]** is switched off in the **[Factory Maintenance>>]** menu.

3.13.4 Software Version

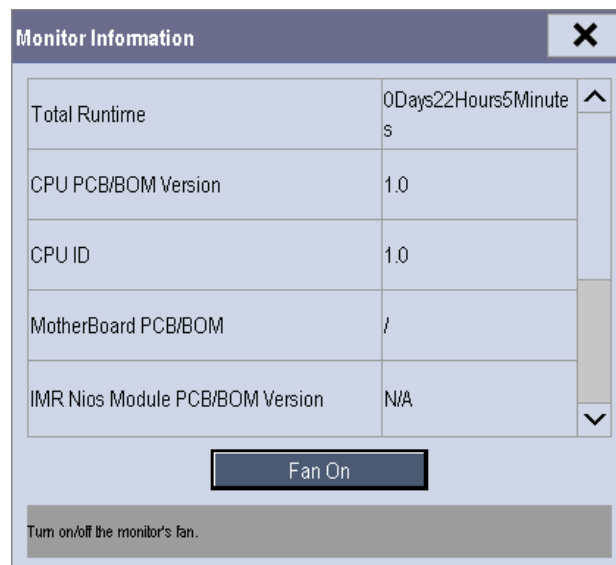
Selecting [**Software Version**] will show software version information. The [**Software Version**] menu is as follows:

A screenshot of a 'Software Version' dialog box. It has a title bar with the text 'Software Version' and a close button (X). The main area contains a table with two columns: the component name and its version. The components listed are System Software Version, Main Power Board, Boot Module, Keypad Module, Recorder Module, and MotherBoard FPGA Logic/Nios Software Version. A vertical scrollbar is on the right side of the table. Below the table is a grey rectangular area.

System Software Version	02.09.00
Main Power Board	1.0
Boot Module	1.0
Keypad Module	1.0
Recorder Module	N/A
MotherBoard FPGA Logic/Nios Software Version	0f.001

3.13.5 Monitor Information

Selecting [**Monitor Information**] will show the status of the patient monitor. Monitor information is displayed as follows:

A screenshot of a 'Monitor Information' dialog box. It has a title bar with the text 'Monitor Information' and a close button (X). The main area contains a table with two columns: the status name and its value. The statuses listed are Total Runtime, CPU PCB/BOM Version, CPU ID, MotherBoard PCB/BOM, and IMR Nios Module PCB/BOM Version. A vertical scrollbar is on the right side of the table. Below the table is a 'Fan On' button and a grey rectangular area with the text 'Turn on/off the monitor's fan.'

Total Runtime	0Days22Hours5Minutes
CPU PCB/BOM Version	1.0
CPU ID	1.0
MotherBoard PCB/BOM	/
IMR Nios Module PCB/BOM Version	N/A

Fan On

Turn on/off the monitor's fan.

(See the above sections for detailed test procedures and contents)

3-46

ECG performance test		
ECG waves are displayed correctly without noise and the HR value is within 80 ± 1 bpm.		
ECG Lead Off alarm behaves correctly.		
Paced signals are detected and pace pulse marks are displayed when [Paced] is set to [Yes]		
The difference between the amplitude of the ECG calibration square wave and that of the wave scale is not greater than 5%.		
Resp test		
The Resp wave is not distorted and the Resp value is within 40 ± 2 rpm.		
SpO₂ test		
Measure SpO ₂ on a healthy person's finger and a Pleth wave and PR value are displayed. The displayed SpO ₂ value is within 95%-100%		
NIBP test		
The difference is within ± 3 mm when 0, 50 or 200 mmHg is set for NIBP accuracy test.		
There is no leakage with NIBP, or the manual leakage test result does not exceed 6mmHg/min.		
Temp test		
The value displayed for each Temp channel of the monitor is within $37 \pm 0.1^\circ\text{C}$.		
IBP test		
The static pressure value displayed for each IBP channel is within 200 ± 2 mmHg.		
The ART and LV waves for each IBP channel are displayed correctly.		
C.O. test		
The TB value displayed on the monitor is within $37 \pm 0.1^\circ\text{C}$.		
The displayed C.O. value is within $5 \pm 0.25\text{L/min}$.		
Mainstream CO₂ test		
The mainstream CO ₂ is zeroed successfully and the waveform baseline recovers to zero.		
CO ₂ Apnea alarm behaves correctly.		
The displayed CO ₂ value is within $6 \pm 0.05\%$.		

Sidestream CO₂ test		
Block the gas inlet of the module or watertrap. The sidestream CO ₂ flowrate is slower than 10ml/min and an alarm of CO ₂ Filterline Err is given. It indicates that there is no leakage.		
The displayed CO ₂ value is within $6 \pm 0.05\%$.		
Microstream CO₂ test		
Block the gas inlet of the module or watertrap. An alarm of CO ₂ Filterline Err is given. It indicates that there is no leakage.		
The displayed CO ₂ value is within $6 \pm 0.05\%$		
AG test		
When AG flowrate is slower than 10ml/min, an alarm of AG Airway Occluded is given. It indicates that there is no leakage.		
The fan inside the AG module works properly.		
The measurement accuracy of CO ₂ , N ₂ O, O ₂ and AA (AA represents an anaesthetic agent) meets the product specifications in the Operator's Manual.		
ICG test		
The measured results are as follows: HR=60±2 bpm, VI=61±4/1000s, TFC=32±2/kOhms.		
The measured results are as follows: HR=70±2 bpm, VI=48±4 /1000s, TFC=32±2 / kOhms.		
BIS test (you can select either method to perform the test)		
Method 1: The BIS value measured on healthy, wide-awake adult is within 80-100.		
Method 2: Connect to the BIS simulator to perform a cyclic impedance check. The EEG wave and BIS numeric are displayed on the monitor.		
RM test		
The displayed TV is within 500±50ml and RR is within 20±1rpm.		
CCO/SvO₂ test		
The CCO/SvO ₂ numerics displayed on the patient monitor and Vigilance monitor are consistent.		
The waves (at the ECG signal output port) displayed on the oscillograph are consistent with the ECG calibration waves displayed on the monitor screen.		

The amplitude of electrical level at the signal output port of MAP, CVP and SpO ₂ are 5±0.25V, 5±0.25V and 10±0.5V respectively.		
PiCCO test		
The detected catheter type accords with the setting of the Pulsion Calbox, and the measurement errors of TB and TI are within ±0.1℃.		
The displayed static pressure values of pArt and pCVP are no more than (200±2) mmHg.		
The waveforms of pArt and pCVP are displayed correctly.		
ScvO₂ test		
The accuracy of ScvO ₂ measurements is (50±3)% and (80±3)%.		
NMT test		
The measurement of TOF-Ratio is between 79 and 81.		
The message “Test passed. The function of NMT sensor is OK” is presented if sensor check completes successfully.		
EEG test		
The result of sensor check is pass, and EEG measurements are displayed on the monitor.		
Nurse call relay performance test		
The relay contacts are close when an alarm occurs.		
Analog output performance test		
The waves displayed on the oscillograph are identical with those displayed on the monitor.		
Electrical safety tests		
Refer to <i>A Electrical Safety Inspection</i> . All the electrical safety tests should be passed.		
Touchscreen calibration		
The touchscreen is calibrated successfully.		
Recorder check		
The recorder can print ECG waves correctly and the printout is clear.		

Set the recorder to some problems such as out of paper, paper jam, etc. the monitor gives corresponding prompt messages. After the problem is removed, the recorder is able to work correctly.		
Automatic alarm recording for each parameter functions correctly when parameter alarms occur.		
Network print test		
The network printer can print out ECG reports correctly.		
Device integration check		
[Devices Integrated] window can display the type of the external device, ventilation mode, and corresponding parameters normally.		
Battery check		
The monitor can operates correctly from battery power when an AC power failure accidentally occurs.		
T5 patient monitor can operate independently on a single battery.		
The operating time of the battery meets the product specification.		

4 Troubleshooting

4.1 Introduction

In this chapter, patient monitor problems are listed along with possible causes and recommended corrective actions. Refer to the tables to check the patient monitor, identify and eliminate the troubles.

The troubles we list here are frequently arisen difficulties and the actions we recommend can correct most problems, but not all of them. For more information on troubleshooting, contact our Customer Service Department.

4.2 Part Replacement

Printed circuit boards (PCBs), major parts and components in the patient monitor are replaceable. Once you isolate a PCB you suspect defective, follow the instructions in *Repair and Disassembly* to replace the PCB with a known good one and check that the trouble disappears or the patient monitor passes all performance tests. If the trouble remains, exchange the replacement PCB with the original suspicious PCB and continue troubleshooting as directed in this chapter. Defective PCB can be sent to us for repair.

To obtain information on replacement parts or order them, refer to *Parts*.

4.3 Patient Monitor Status Check

Some troubleshooting tasks may require you to identify the hardware version and status of your patient monitor.

1. To view the information on system start time, self check, etc., select [**Main Menu**]→[**Maintenance >>**]→[**Monitor Information >>**].
2. You can also view the information on the monitor's current status by selecting [**Main Menu**]→[**Maintenance>>**]→[**Factory Maintenance>>**]→enter the required password →[**Monitor Information >>**].

4.4 Software Version Check

Some troubleshooting tasks may require you to identify the configuration and software version of your patient monitor.

1. To view information on the system configuration and system software version, Select [Main Menu]→[Maintenance>>]→[Software Version>>].
2. You can also view the information on system software version and module software version by selecting [Main Menu]→[Maintenance>>]→[Factory Maintenance>>]→ enter the required password →[Software Version>>].

4.5 Technical Alarm Check


Before troubleshooting the patient monitor, check for technical alarm message. If an alarm message is presented, eliminate the technical alarm first. For detailed information on technical alarm message, possible cause and corrective action, refer to the patient monitor's Operation Manual.

4.6 Troubleshooting Guide

4.6.1 Power On/Off Failures

Symptoms	Possible Cause	Corrective Action
The patient monitor fails to start. AC LED or battery LED does not light	AC mains not connected or battery too low	Check that AC mains is properly connected or battery capacity is sufficient.
	Power supply protection	Refer to 4.6.9 Power Supply Failures .
	Cables defective or poorly connected	1. Check that the cables from power switch & LED board to button board, button board to main board, and power module to main board are correctly connected. 2. Check that cables and connectors are not damaged.
	Power switch & LED board defective	Replace the power switch & LED board.
	Power module defective	Replace the power module.
	Mother board Defective	Replace the mother board.

4.6.2 Display Failures

Symptoms	Possible Cause	Corrective Action
Integrated display is blank but the patient monitor still works correctly.	Cables defective or poorly connected.	1. Check that the cable from the display to the mother board and the cables from the backlight board respectively to the button board and the display are correctly connected. 2. Check that the cables and connectors are not damaged.
	Backlight board defective	Replace the backlight board.
	Power module defective	Replace the power module.
	Display defective	Replace the display.
Secondary display does not function.	Cables defective or poorly connected.	1. Check that the cable between the display and the patient monitor is correctly connected. 2. Check that the cables and connectors are not damaged.
	DVI interface board defective	Replace the DVI interface board.
Secondary display displays snows or flashing specks	Cables defective or poorly connected.	1. Check that the cable between the display and the patient monitor is correctly connected. 2. Check that the cables and connectors are not damaged.
	DVI interface board defective	Replace the DVI interface board.
	The mother board is damaged.	Replace the mother board.
Images overlapped or distorted	FPGA error.	Update or upgrade FPGA.
	Cables defective or poorly connected.	1. Check that the cable between the display and mother board is correctly connected. 2. Check that the cables and connectors are not damaged.
Touchscreen does not response	Touchscreen disabled	Check if there is a symbol  shown above the [Main Menu] QuickKey. If yes, press the [Main Menu] QuickKey for more than 3s to enable the touchscreen.

	Cables defective or poorly connected.	<ol style="list-style-type: none"> 1. Check that the cables from the touchscreen to the touchscreen control board, the touchscreen control board to the button board, and the button board to the mother board are correctly connected. 2. Check that the cables and connectors are properly connected
	Touchscreen control board defective	Replace the touchscreen control board
	Button board defective.	Replace the button board.
	Touchscreen defective.	Replace the touchscreen
	Mother board defective	Replace the mother board
Touch position invalid	Touchscreen not calibrated	Calibrate the touchscreen

4.6.3 Module Rack Failures

Symptoms	Possible Cause	Corrective Action
SMR		
SMR cannot identify parameter modules	Extension Cable defective or poorly connected	<ol style="list-style-type: none"> 1. Check that the cable between SMR and main unit is properly connected 2. Check that the connecting cables and connectors are not damaged. 3. Check that contact screws on SMR are tightly screwed and properly contact the SMR.
	Defective parameter module	Replace the suspicious parameter module with a known good module. Check if the patient monitor identifies the replacement module. If yes, it means that the original one is defective.
	Wrong communication board software revision	Upgrade the program of the module or SMR.
	Module (in some slots) not recognized	Replace the Nios II module. Replace the 8-slot module rack communication board.

	Power supply failure	<ol style="list-style-type: none"> 1. Check if the voltage between two contact screws in any slot reaches 12V DC. If yes and the parameter module functions properly and the PCB assembly in SRM might fail. 2. If there is no 12 VDC power sent to the SMR, check whether the power voltage output to the USB_Hub board by the power module reaches 12V. If yes, the fuse of the USB interface board might blow. Replace the USB_Hub board.
	Cable defective or poorly connected	<ol style="list-style-type: none"> 1. Check that the cable between SMR interface board and communication board is properly connected. 2. Check that connecting cables and connectors are not damaged.
	Nios II module loose or failure	<ol style="list-style-type: none"> 1. Check that Nios II module is correctly plug ed 2. If the symptom persists, replace the Nios II module.
	SMR interface board failure	Replace the SMR interface board.
	SMR communication board failure	Replace the SMR communication board.
	USB_Hub board failure	Replace the USB_Hub board.
	Mother board failure	Replace the mother board.
Integral module rack		
Integral module rack cannot identify parameter modules	Module failure	Replace parameter module. If a new module is identified, the original one is defective.
	Cable defective or poorly connected	<ol style="list-style-type: none"> 1. Check that the cables from 3-slot module rack communication board to MPM module rack communication board, module rack to mother board are properly connected. 2. Check that connecting cables and connectors are not damaged.
	Wrong communication board software revision	Upgrade the program of the module or Integral module rack.
	Module (in some slots) unrecognized	Replace the corresponding module rack communication board.

	Power supply to integral module rack abnormal	<ol style="list-style-type: none"> 1. Check if voltage between two contact screws in any slot reaches 12VDC. If yes and the parameter module functions, PCB assembly in the SMR might fail. 2. If there is no 12V sent to the integrated module rack, check that power module output voltage to mother board reaches 12V DC. If yes, mother board might fail.
	3-slot or MPM module rack communication board failure	Replace the 3-slot or MPM module rack communication board.
	Nios II module failure	Replace the Nios II module.
	Mother board failure	Replace the mother board.

4.6.4 Alarm Problems

Symptoms	Possible Cause	Corrective Action
The alarm lamp is not light or extinguished but alarm sound is issued	Cable defective or poorly connected	<ol style="list-style-type: none"> 1. Check that cables from alarm LED board to button board and button board to mother board are properly connected. 2. Check that connecting cables and connectors are not damaged.
	Alarm LED board failure	Replace the alarm LED board.
	Button board failure	Replace the button board.
	Mother board failure	Replace the mother board.
No alarm sound is issued but alarm lamp is light	Audio alarm disabled	Select [Main Menu]→[Maintenance >>]→[User Maintenance >>]→enter the required password→[Alarm Setup >>], and then in the popup menu, set [Minimum Alarm VolumeAlm Sound] to appropriate setting.[On] In the [Others] window of the [Alarm Setup] menu, set [Alm Volume] to appropriate setting.
	Cable defective or poorly connected	<ol style="list-style-type: none"> 1. Check that cable between speaker and mother board is properly connected. 2. Check that connecting cables and connectors are not damaged.
	FPGA audio logic error	Upgrade the audio logic part of the FPGA program.
	Speaker failure	Replace the speaker.
	Mother board failure	Replace the mother board.

4.6.5 Button and Knob Failures

Symptoms	Possible Cause	Corrective Action
Buttons do not work	Cable defective or poorly connected	1. Check that cable between button board and mother board is properly connected. 2. Check that connecting cables and connectors are not damaged.
	Button board failure	Replace button board.
Knob does not work	Cable defective or poorly connected	1. Check that cables from knob to button board, and button board to mother board are properly connected 2. Check that connecting cables and connectors are undamaged.
	Knob failure	Replace the knob encoder.
	Button board failure	Replace the button board

4.6.6 Recorder Failures

Symptoms	Possible Cause	Corrective Action
No printout	Recorder module disabled	1. Check if the recorder status LED lights 2. If yes, enable the module in [Factory Maintenance] menu. Otherwise, check for other possible causes.
	Paper reversed	Re-install the paper roll.
	Cable defective or poorly connected	1. Check that cable between recorder and mother board is properly connected. 2. Check that connecting cables and connectors are not damaged.
	Recorder power supply failure	Check if the power module outputs 5 V DC and 12V DC correctly.
	Recorder failure	Replace the recorder.
Poor print quality or paper not feeding properly	Paper roll not properly installed	Stop the recorder and re-install the paper roll.
	Print head dirty	1. Check the thermal print head and the paper roller for foreign matter. 2. Clean the thermal print head with an appropriate clean solution.
	Print head failure	Replace the print head.
	Recorder failure	Replace recorder.

4.6.7 Output Interface Failures

Symptoms	Possible Cause	Corrective Action
No analog signals or nurse call signals are issued	Respective output disabled	1. Select [Main Menu]→[Analog Output Setup]→set [Analog Output] to [On].
	USB_Hub board cable loose	1. Check that cable between USB_Hub board and mother board is properly connected. 2. Check that connecting cables and connectors are not damaged.
	USB_Hub board failure	Replace the USB_Hub board.
	Mother board failure	Replace the mother board.
Device with USB port does not function (Assume that the peripheral devices are good)	Cable defective or poorly connected	1. Check that cable between USB_Hub board and mother board is properly connected. 2. Check that connecting cables and connectors are not damaged.
	USB_Hub board failure	Replace the USB_Hub board.
	Mother board failure	Replace the mother board.

4.6.8 CF Card Problems

Symptoms	Possible Cause	Corrective Action
CF card malfunctions	Wrong CF card or small memory space	Use only SanDisk-manufactured CF storage cards. Those with 1GB memory space are recommended.
	CF card full; data error; CF card error	Format CF card on PC.
	CF card failure	Replace the CF card.
	Cable defective or poorly connected	1. Check that the cable between CF card board and mother board is correctly connected. 2. Check that connecting cables and connectors not damaged.
	CF card board failure	Replace the CF card board.
	Mother board failure	Replace the mother board.

4.6.9 Power Supply Failures

Symptoms	Possible Cause	Corrective Action
Different battery voltages	Battery failure	Replace battery.
	Cable defective or poorly connected	1. Check that the cable between battery interface board and power module is correctly connected. 2. Check that cables and connectors are not damaged.
	Power board failure	Replace the power board.
Battery capacity is too low	Battery failure	Replace battery.
	Cable defective or poorly connected	1. Check that the cable is correctly connected. 2. Check that connecting cables and connectors are not damaged.
	Power board failure	Replace the power board.
Battery cannot be recharged	Battery failure	Replace battery and recharge the replacement battery. If the replacement battery can be recharged, the original one fails.
	Cable defective or poorly connected	1. Check that cable between battery interface board and power module is correctly connected. 2. Check that cables and connectors are not damaged.
	Power board failure	Replace power board
No +3.3 V output	1. Power supply protected 2. Power board failure	1. Turn off the patient monitor then restart it. 2. If the problem remains, disconnect the AC mains for 5 s and reconnect it, then restart the patient monitor. 3. If the problem still remains, replace the power board.
No +5.0 V output		
No +12 V output		

NOTE

- When the power module has a failure, it may cause problems to other components, e.g. the monitor suddenly breaks down during start-up, as the power module may have a power supply protection. In this case, troubleshoot the power module per the procedure described in the table above.
 - Components of the main unit, SMR and parameter modules are powered by the power module. In the event that a component malfunctions, check if the operating voltage is correct. Refer to *2 Theory of Operation* for the operating voltage and measurement points of each component.
-

4.6.10 Network Related Problems

Symptoms	Possible Cause	Corrective Action
Frequent dropouts and network disconnects	Incorrect LAN cable connection	Check LAN cable connection. LAN cable shall not be longer than 50 m.
	Incorrect IP address configuration	Check for IP address conflict. Reconfigure IP address.
The patient monitor is connected to a LAN but cannot view other patients in the View Others mode	Incorrect LAN cable connection	Check LAN cable connection. LAN cable shall not be longer than 50m.
	Excessive requests for viewing the patient monitor at the same time	A patient monitor can only be viewed by 4 other patient monitors at the same time under the View Others mode. The excessive view requests system will be ignored.
	Incorrect IP configuration	Check for IP address conflict. Reconfigure IP address.
	USB_Hub board failure	Replace the USB_Hub board.

4.6.11 Software Upgrade Problems

Symptoms	Possible Cause	Corrective Action
Bootstrap upgrade fails	Power failure or unintended power off during bootstrap upgrade	Return the CPU board to factory for repair.
Program upgrade fails	Incorrect network connection	1. Check that network connector, not iView connector, on the patient monitor is used. 2. Make sure that the hub or switch run normally. Check that net twines are of the right type and have been connected correctly.
	Wrong upgrade package has been downloaded	Upgrade package shall be .pkg files. Select package according to system requirement.
	Incorrect IP address configuration	Configure a fixed IP address in range C as specified for the patient monitor. We recommend not to upgrade a program when the patient monitor is connected to a network with multiple PCs.

4.6.12 Technical Alarm Messages

Please refer to the Operator's manual.

4.6.13 M51A Self Test Information

New MPM module applies the integrative parameter board (ECG ASIC).

MPM Selftest Item	Test Value	Test Value (New MPM module)	Corrective Action
DSP selftest information	Not F	Not FF	Replace the module
7024 selftest information	Not 7F		
2131 selftest information	Not 1F		
ECG module selftest information	Normal value: 7 for 3/5 lead module; FF for 12-lead module		

4.6.14 Device Integration Failures

Symptoms	Possible Cause	Corrective Action
The [Devices Integrated] window displays nothing after connection.	The ID adapter is not compatible with the external device.	1. Replace the ID adapter. 2. Upgrade the ID adapter in [Factory Maintenance] menu to make the ID adapter match the corresponding external device. See <i>3.II.1 Device Connection and Setup</i> for more about the setup of the ID.
	The serial port adapting cable is not compatible with the external device.	Replace the serial port adapting cable.
	Wrong software version or wrong protocol version of the external device.	Make sure the protocol version and software version are supported by the BeneLink module.
Generate the alarm: [BeneLink Comm Stop].	The application of the BeneLink module is damaged.	Update or upgrade the application of the BeneLink module with the network upgrading tool.
The patient monitor has no response when loading the ID adapter.	The application of the BeneLink module is damaged.	Update or upgrade the application of the BeneLink module with the network upgrading tool.
	The kernel or the document system of the BeneLink module is damaged.	Return the BeneLink module to factory for repair.

5 Repair and Disassembly

5.1 Tools

During disassembly and replacing, the following tools may be required:

- Phillips screwdrivers
- Small flat-bladed screwdrivers
- Contact spanner
- Tweezers
- Sharp nose pliers
- Sleeve

5.2 Preparations for Disassembly

Before disassembling the monitor, finish the following preparations:

- Stop monitoring the patient, turn off the monitor and disconnect all the accessories and peripheral devices.
- Disconnect the AC power source and take out both of the batteries.
- Pull off all the modules in the integral module rack. If the SMR is connected, disconnect the SMR from the monitor and then remove all the modules in it.



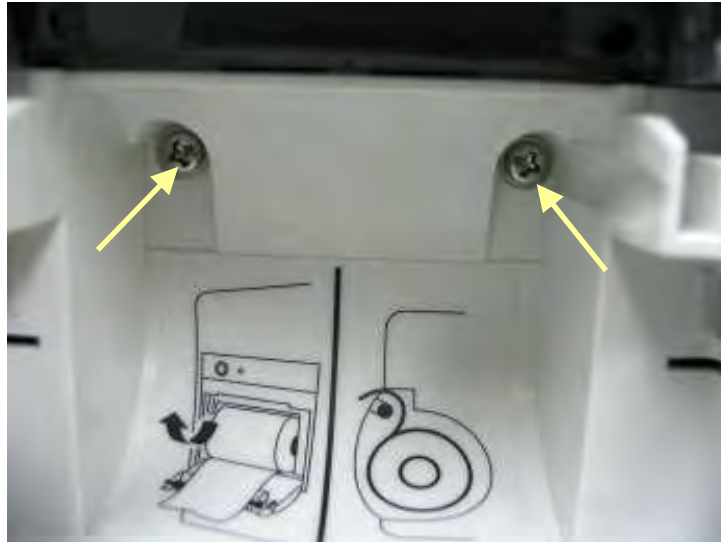
WARNING

- **Before disassembling the monitor, be sure to eliminate the static charges first. When disassembling the parts labeled with static-sensitive symbols, make sure you are wearing electrostatic discharge protection such as antistatic wristband or gloves to avoid damaging the equipment.**
 - **Put the cables or wires in place when reassemble the monitor to avoid short circuit.**
 - **When assembling the monitor, be sure to select proper screws. If an unfit screw is tightened by force, the monitor may be damaged and the screw or the part may fall off during use to cause unpredictable damage or human injury.**
 - **Be sure to follow the correct sequence to disassembly the monitor. Otherwise, the monitor may be damaged permanently.**
 - **Be sure to disconnect all the cables before disassembling any parts. Be sure not to damage any cables or connectors.**
 - **Be sure to place the monitor face up when disassembling it. Otherwise, the screen or the knob may be scratched or damaged.**
-

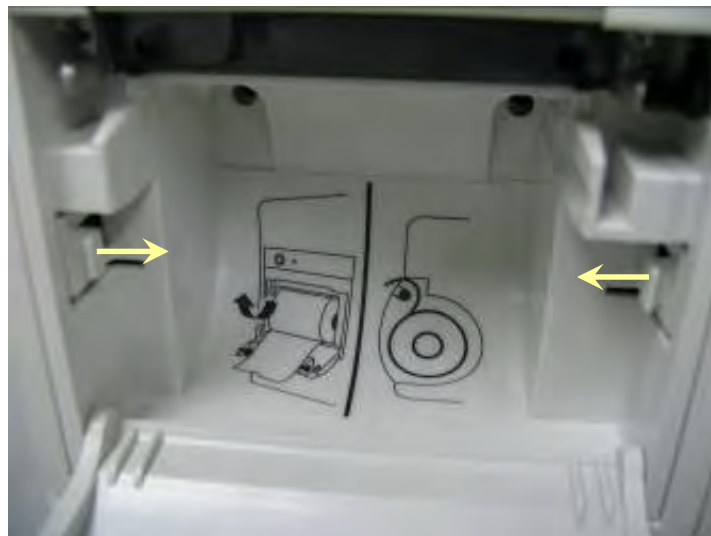
5.3 Disassembling Procedure

5.3.1 Removing the Recorder

1. Open the recorder door and unscrew the two M3×6 screws.



2. Pull the two clips in the directions as indicated and meanwhile pull out the recorder.



NOTE

- Be sure not to damage the connecting cables or connectors when pulling out the recorder.

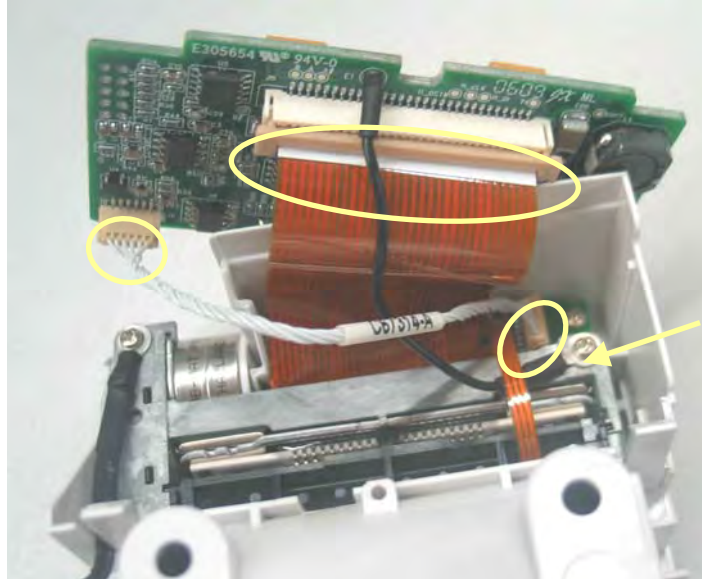
-
3. Unscrew the M3×6 screw and unplug the recorder grounding cable and the cable between the recorder and the mother board.



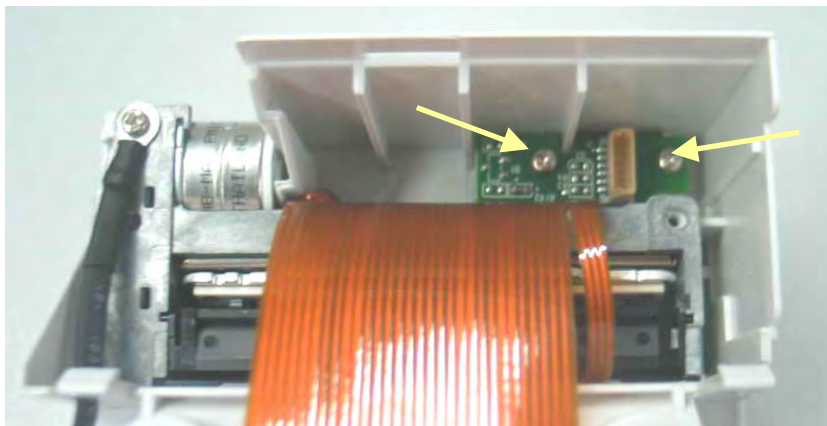
4. Pull the two clips backwards and remove the recorder driving board.



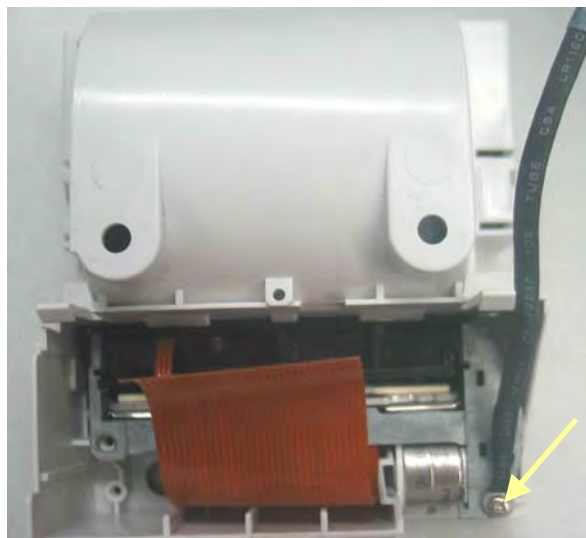
-
5. Pull the press bar upwards about 1 mm and then unplug the flexible cable. Remove the cable that connects the driving board and the button board. Unscrew the PT2×6 screw and remove the drive board's grounding cable. Then take out the recorder driving board.



6. Unscrew the two PT2×6 screws and take out recorder's button board.



-
7. Unscrew the PT2×6 screw. Prize the thermal printhead a little using a small flat-bladed screwdriver to remove the printhead.



5.3.2 Separating the Front and Rear Housing

1. Remove the hook and then unscrew the four M3×12 screws, one of which can be seen only when the battery door is opened.



NOTE

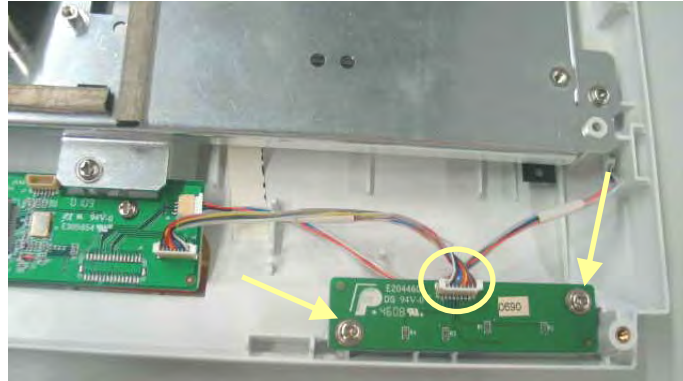
- Exercise care when pulling the front housing out. Be sure not to damage the cables and connectors.
 - Avoid pressing the knob on the table.
-
2. Separate the front and rear housing and unplug the connecting cable between button board and main board as well as the cable between the LCD and the main board.



Avoid pressing the knob on the table.

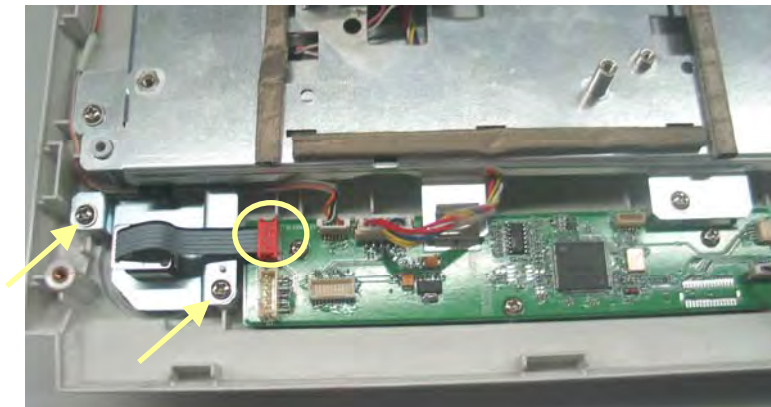
5.3.3 Removing the Power Switch & LED Board

Unplug the connecting cable between power switch & LED board and the button board.
Unscrew the two M3×6 screws and remove the power switch & LED board.



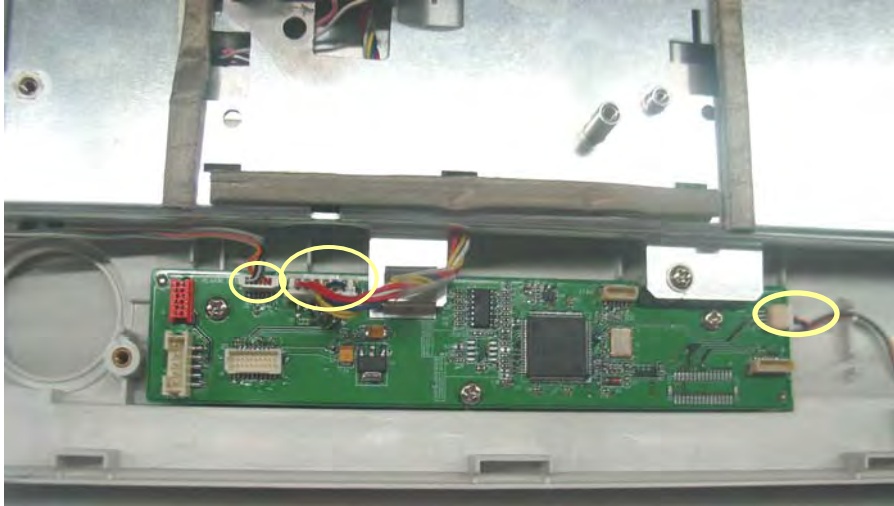
5.3.4 Removing the Knob Encoder

Disconnect the cable that connects the knob encoder and the button board. Then unscrew the two M3×6 screws and take out the knob encoder.

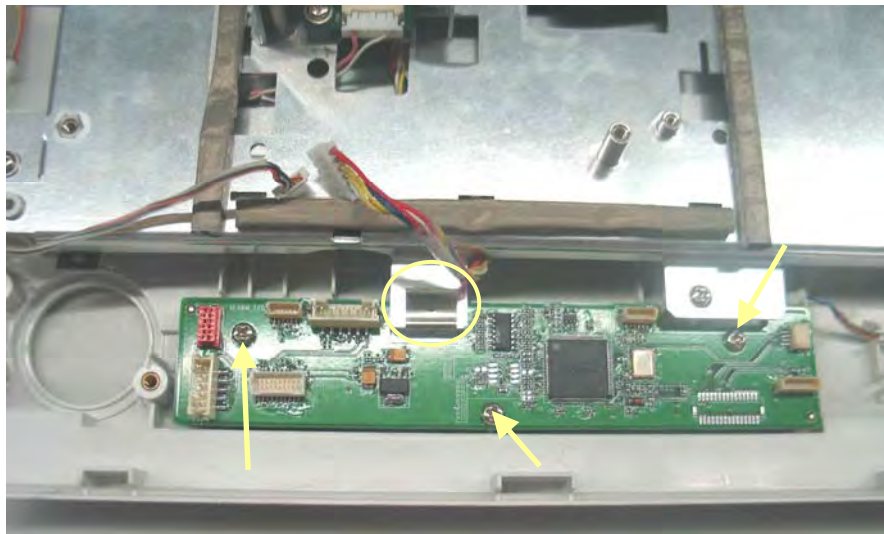


5.3.5 Removing the Button Board

1. Disconnect the cables from the button board to the alarm LED board, backlight board and touchscreen control board.



2. Remove the grounding spring and then unscrew the three PT3×8 screws and take out the button board.

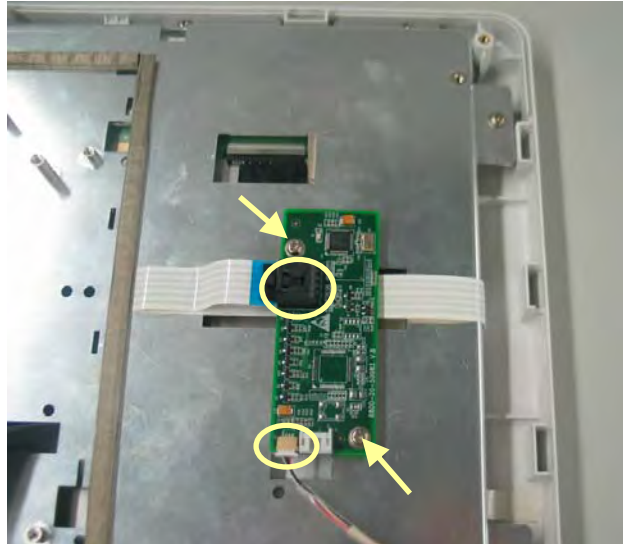


NOTE

- Do not forget the grounding spring when reassembling..

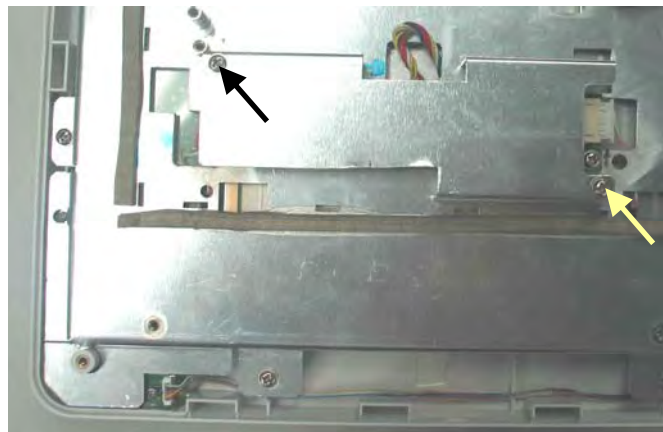
5.3.6 Removing the Touchscreen Control Board

Unplug the cables respectively from the touchscreen and button board to the touchscreen control board. Then, unscrew the two M3×6 screws and remove the touchscreen control board.

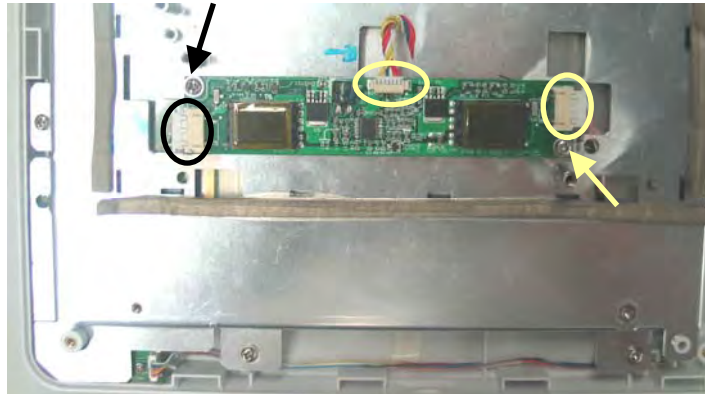


5.3.7 Removing the Inverter

1. Unscrew the two M3×6 screws and remove the backlight board shield.



-
2. Unplug the cables respectively from the button board and the LCD to the backlight board. Then, unscrew the two M3×6 screws to remove the inverter.

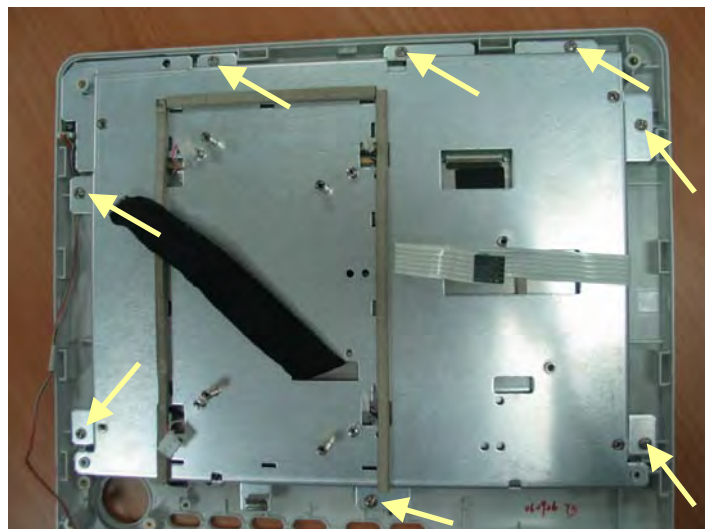


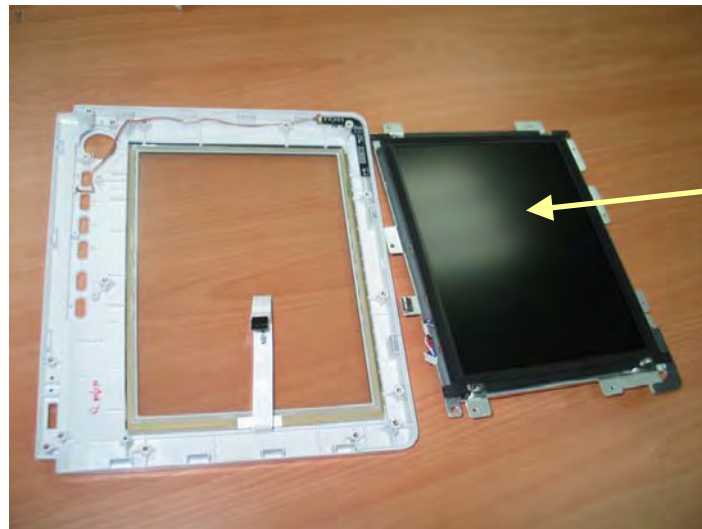
5.3.8 Removing the LCD

CAUTION

- Do not touch the LCD.
 - Disassemble the LCD in an environment as dust-free as possible.
-

1. Unscrew the eight M3×6 screws and remove the LCD with care. Do not touch the LCD surface and prevent it from being contaminated by dust.





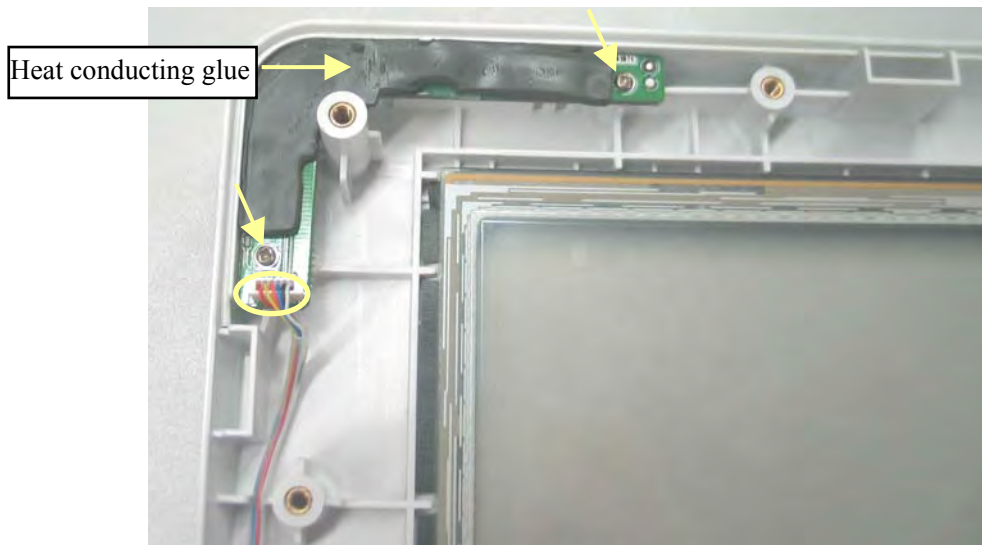
Do not touch the
LCD surface

2. Unscrew the four M3×6 screws underneath the screen cover and then remove the LCD.



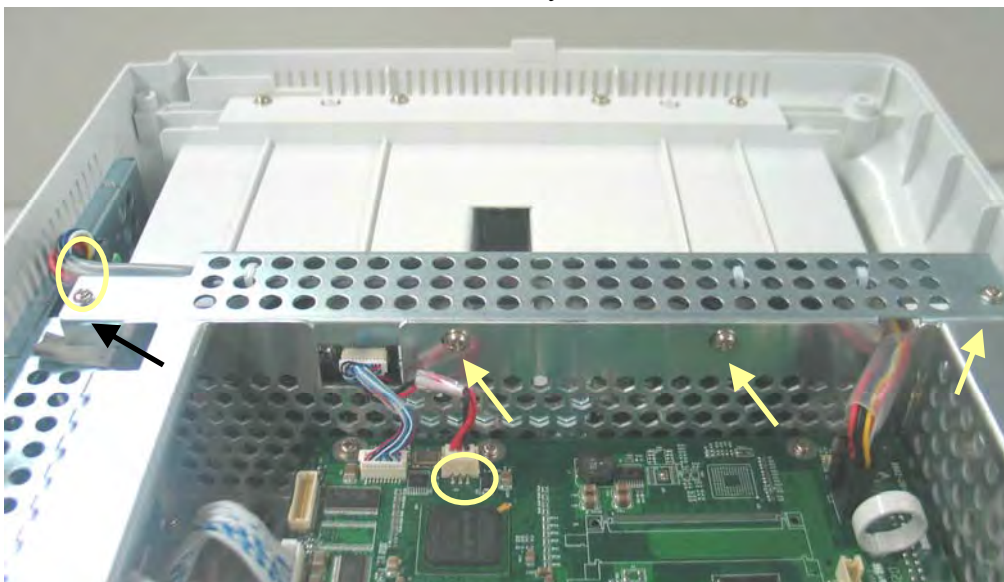
5.3.9 Removing the Alarm LED Board

After removing the LCD, disconnect the cable that connects the alarm LED board and the button board, and then, unscrew the two PT2×6 screws to remove the alarm LED board.



5.3.10 Removing the Fan Assembly

Unplug the cable that connects the fan assembly and the mother board. Then, unscrew the four M3×6 screws and remove the fan assembly.



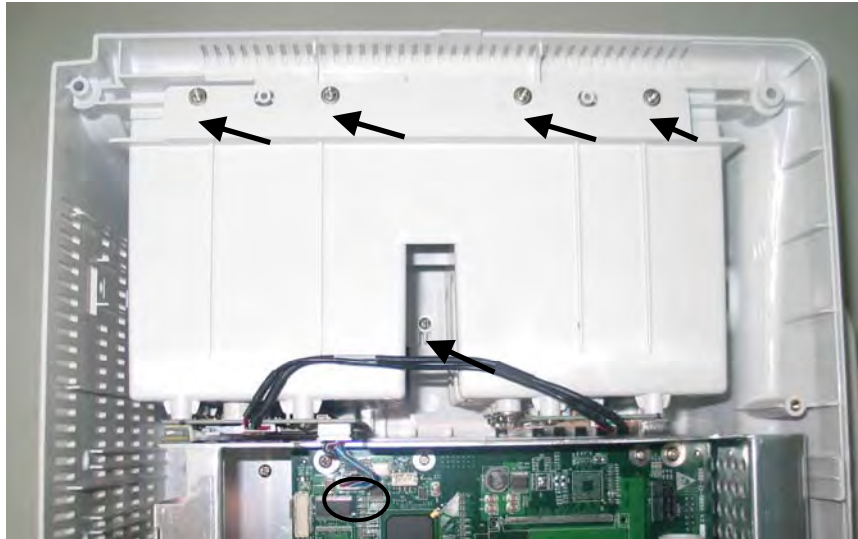
5.3.11 Removing Battery Compartment Assembly

Unplug the cable that connects the battery compartment assembly and the mother board. Then, unscrew the three M3×6 screws and take out the battery compartment assembly.

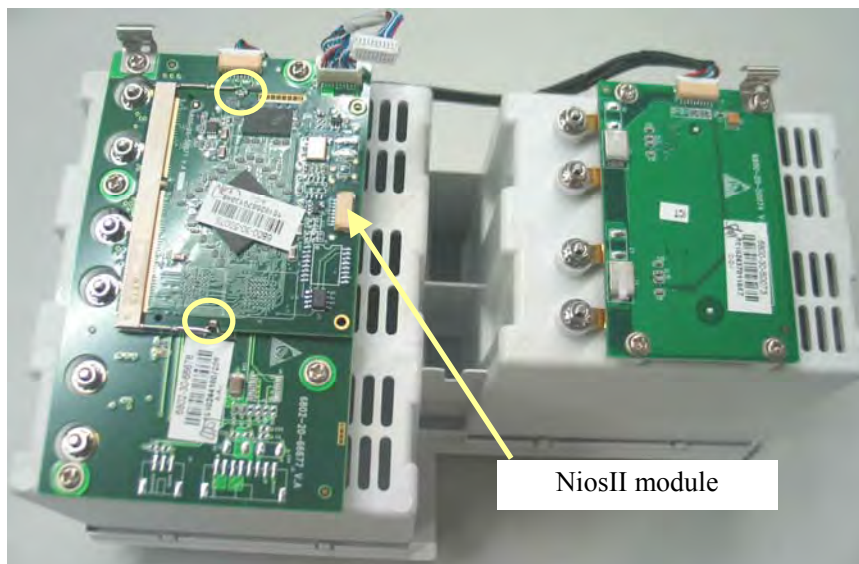


5.3.12 Removing the Integral Module Rack

1. Disconnect the cable that connects the integral module rack and the mother board.
Unscrew the five M3×8 screws and remove the integral module rack.



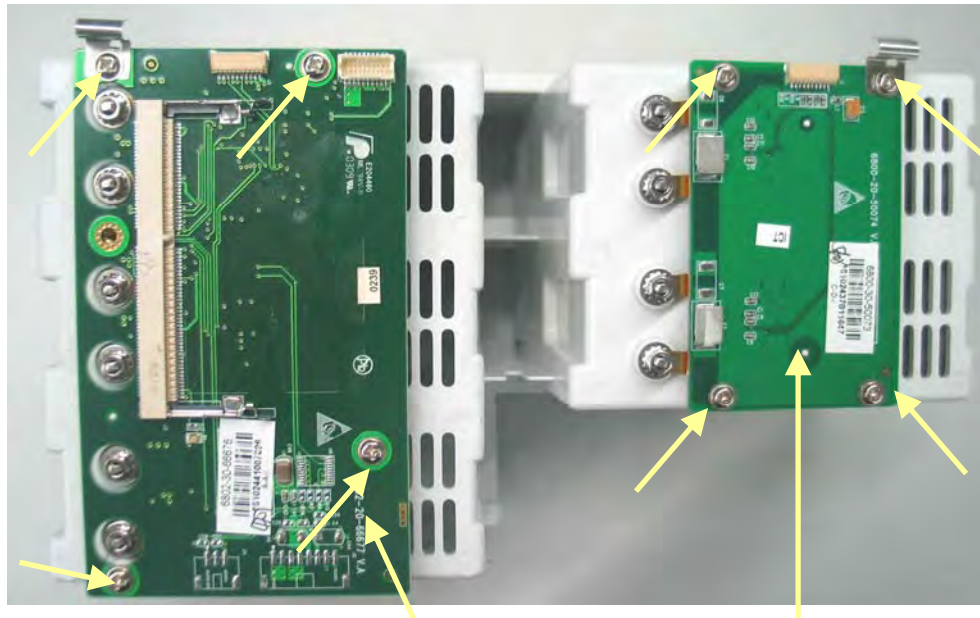
2. Disconnect the cable that connects the MPM module rack communication board and the 3-slot module rack communication board. Release the two snaps and unplug the NiosII module on the 3-slot module rack.



3. Unscrew the eight M3×6 screws and then remove the 3-slot module rack communication board and MPM module rack communication board.

NOTE

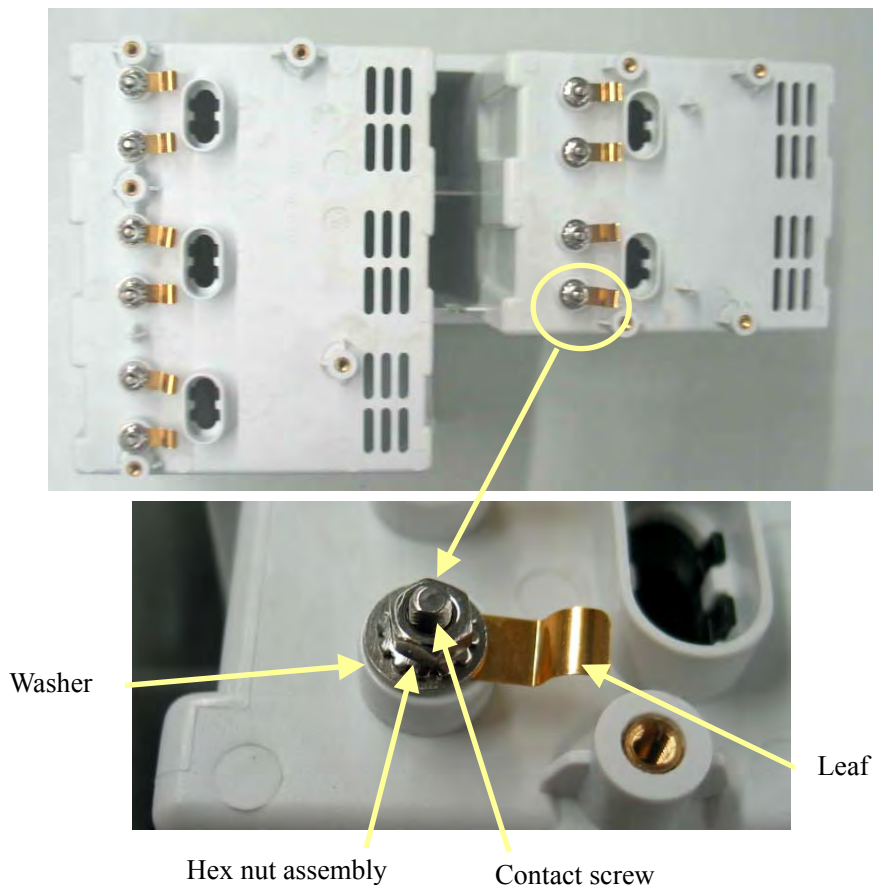
- Do not forget the grounding spring when reassembling..
-



3-slot module rack
communication board

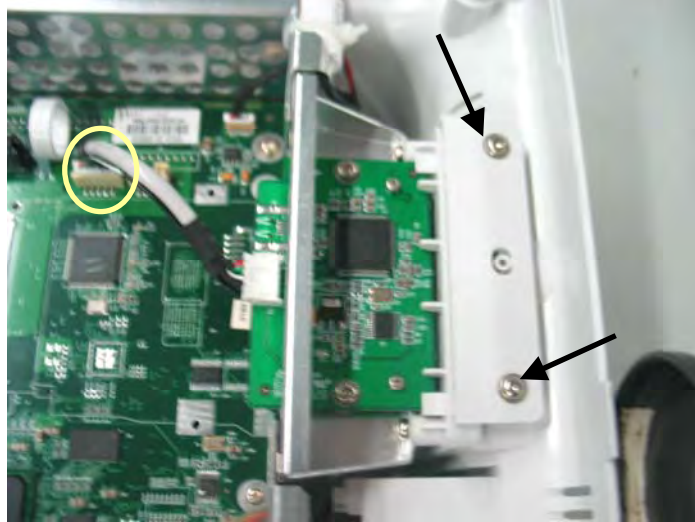
MPM module rack
communication board

-
4. Unscrew the hex nut assy using the sleeve. Then separate the washer, spring and contact screw from each other.



5.3.13 Removing the CF Card Assembly

1. Unplug the cable that connects the CF card assembly and the mother board. Then, unscrew the two M3×6 screws and remove the CF card assembly.



2. Unscrew the two M3×6 screws and remove the CF card assembly with the shield.



NOTE

- When reassembling the CF card assembly, be sure to install the CF card door first.

5.3.14 Removing the wireless AP assembly

5.3.14.1 Removing the internal wireless AP

1. Disconnect the wireless AP cable. Then unscrew the two M3×6 screws to remove the wireless AP assembly.



2. Unscrew the two M3×6 screws to remove the wireless AP.

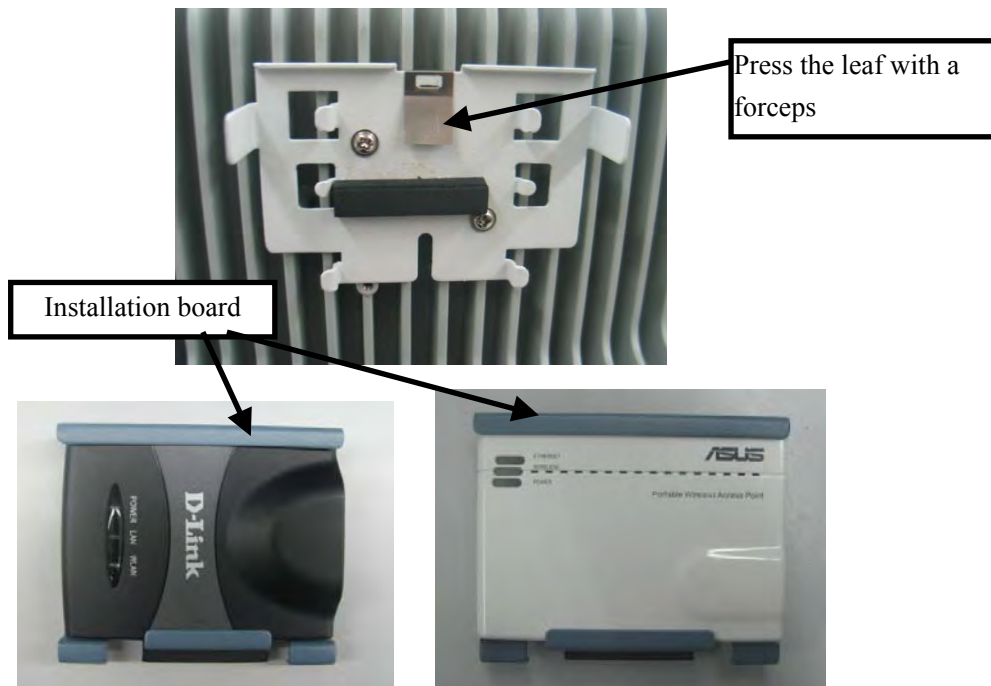


5.3.14.2 Removing the external wireless AP

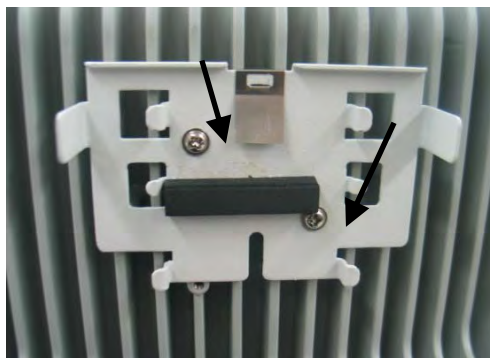
1. Disconnect the network cable and the power cord



2. Press the leaf on the fix board with forceps; push the AP and installation board upwards. Then remove the AP and installation board.



3. Unscrew the two M3×8 screws and take out the fix board.



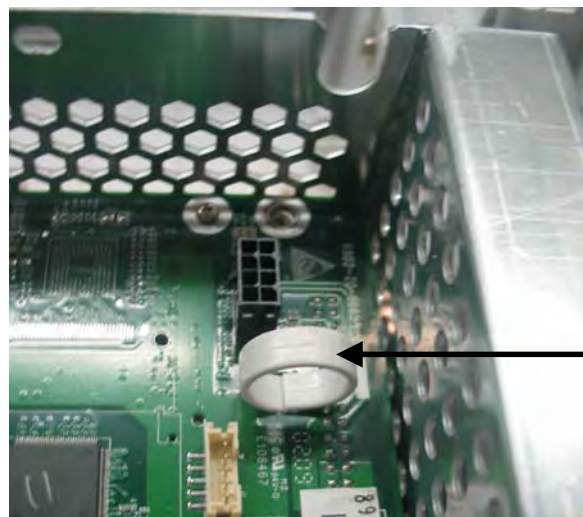
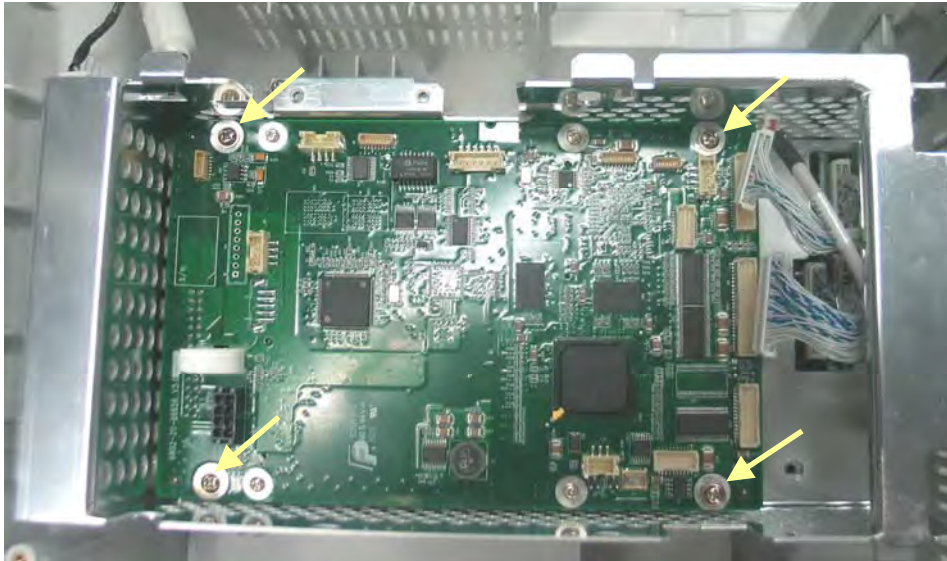
5.3.15 Removing the Main Board

1. Unplug all the cables on the mother board. The numbers beside the connectors indicates what device is connected with the connector.

J4	Button board	J17	speaker
J5	Recorder	J19	LCD
J6	DVI interface board	J9	Wireless AP
J7	Battery interface board	J23	Fan
J10	CF card board	J25	3-slot module rack communication board
J12	power board	J18、J26	USB_Hub board



-
2. Unscrew the 4 M3×6 screws and take out the main board assembly.

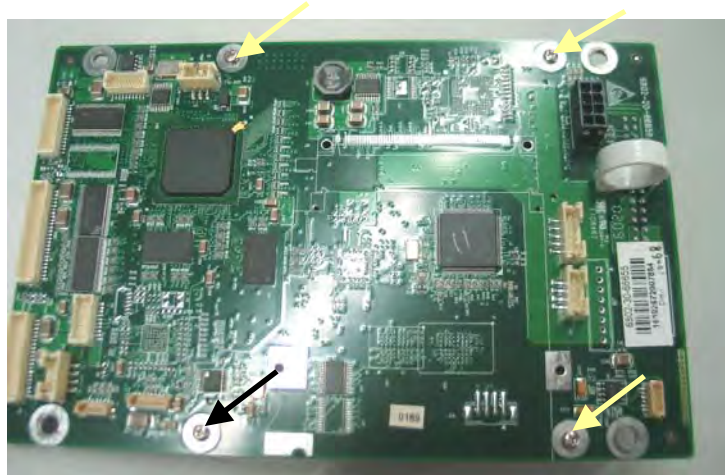


Ring

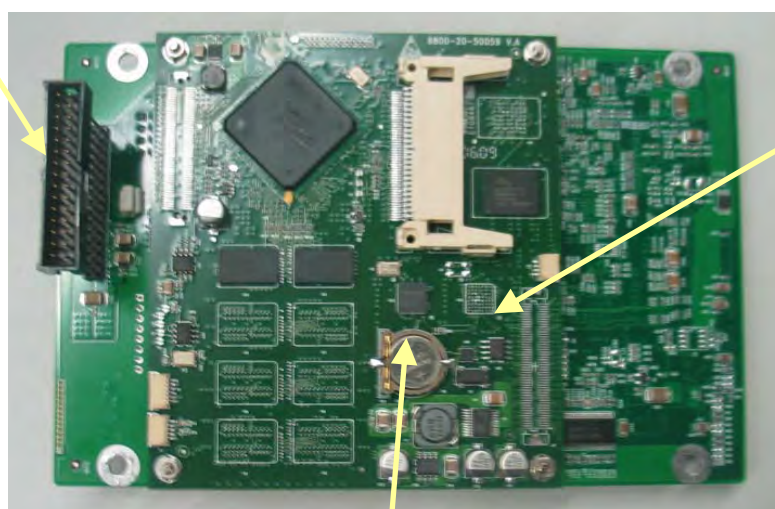
NOTE

- Since the main board assembly is connected with the power module via a butt socket and the CPU radiator may be adhered to it, the main board assembly should be removed with force. Take care not to damage the butt socket.

3. Unscrew the 4 M2.5×6 screws on the CPU board and vertically separate the CPU board and the mother board. Take care not to damage the butt socket.



Butt socket to
power module

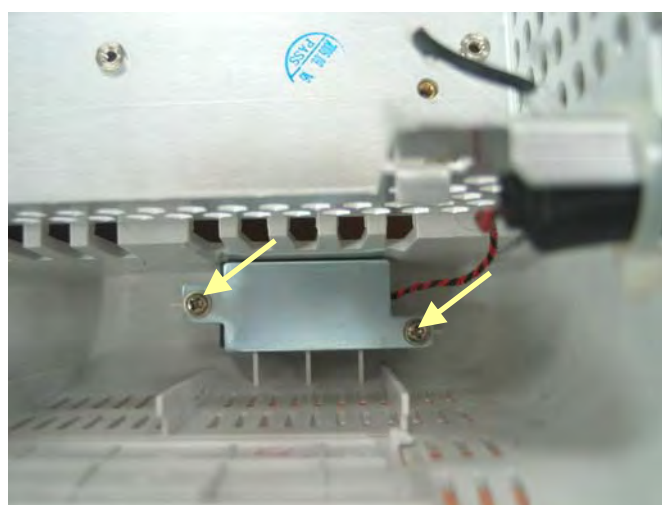


CPU board

Button cell

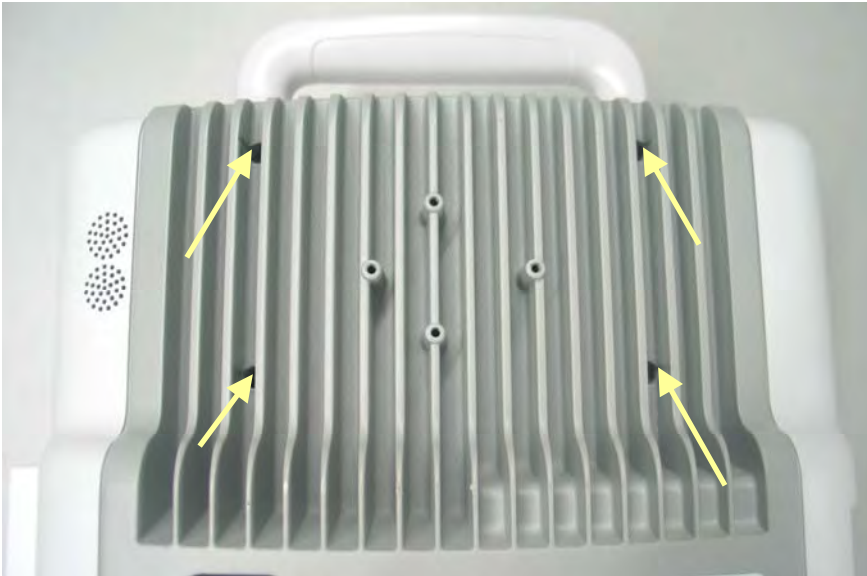
5.3.16 Removing the Speaker

Unscrew the two M3×6 screws and remove the speaker.

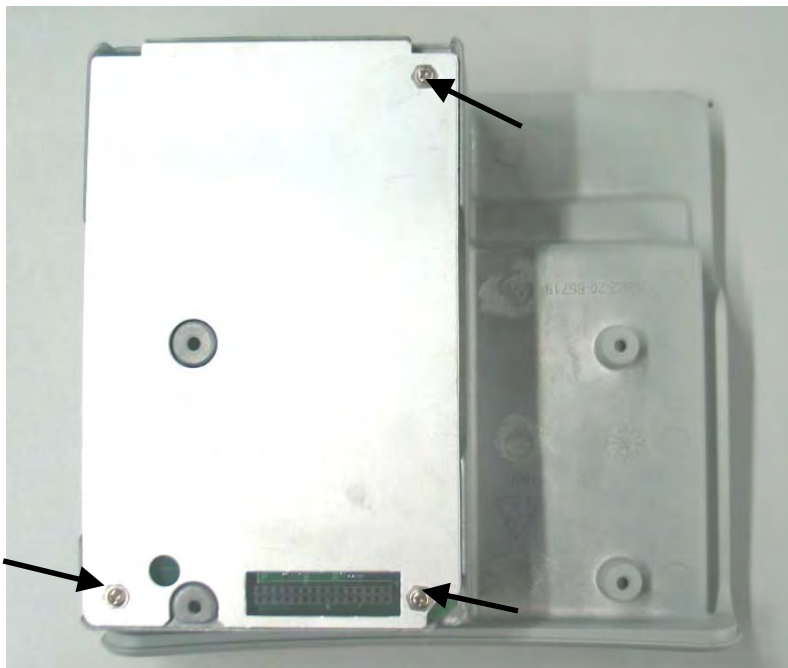


5.3.17 Removing the Power Module Assembly

1. Unscrew the four M3×12 screws and remove the power module assembly.



2. Unscrew the three M3×12 screws and remove the power supply cover.



-
3. Lift the power board with a small flat screwdriver. Then, turn it over.



4. Unplug the cable between the AC input filter and the power board, then remove the power board.



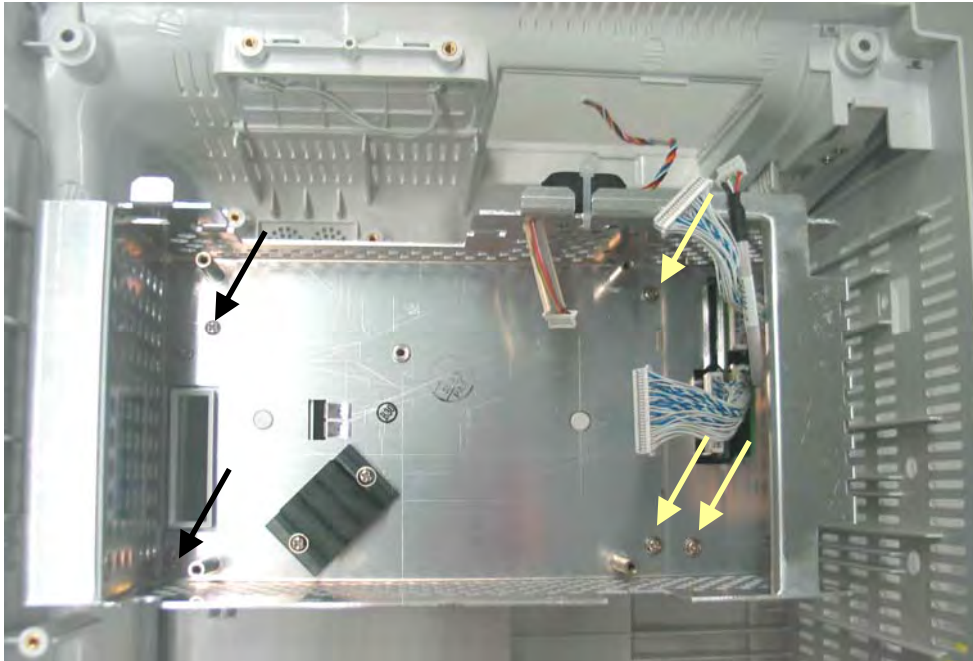
Equipotential Pillar

NOTE

- Since the power board may be adhered to the insulator, be careful not to damage the parts, connectors and cables on the power board when prizing it.

5.3.18 Removing the Main Support

Unscrew the five M3×6 screws and disassemble the main support.

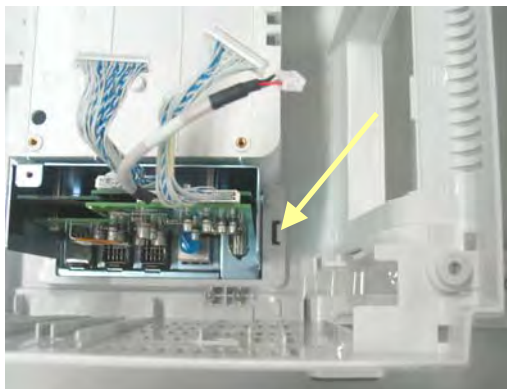


Note

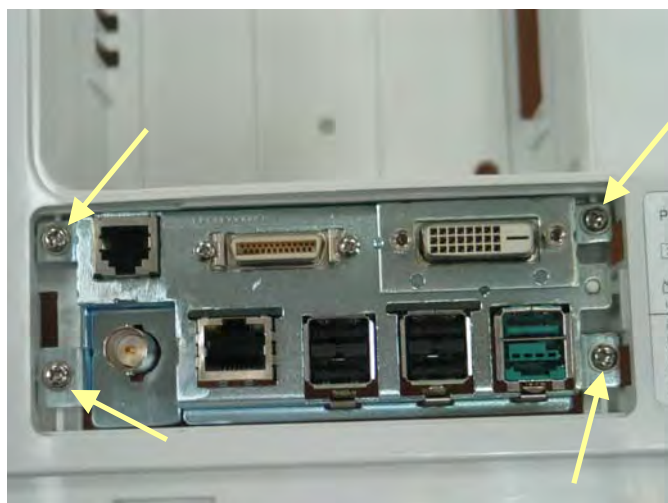
- Be sure to remove the power module first before removing the main support.

5.3.19 Removing the Interface Board Assembly

1. Release the clip that locks the interface board and push it outwards.



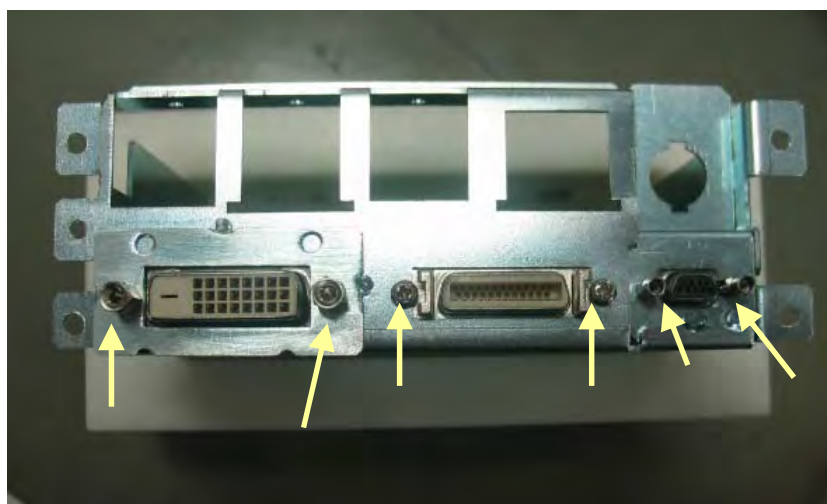
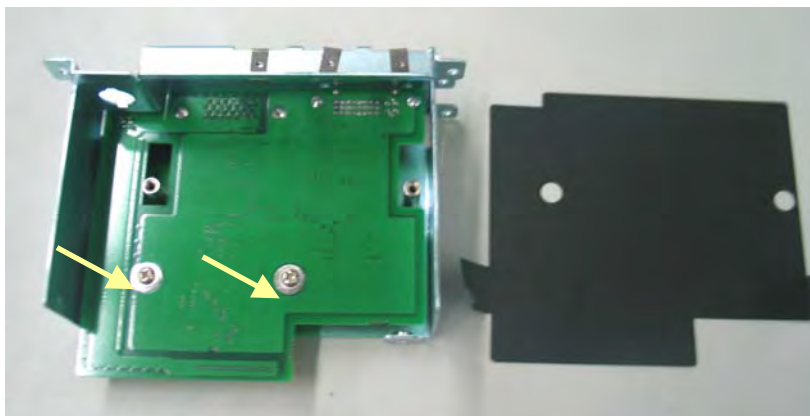
-
2. Open the interface board cover and unscrew the four M3×6 screws, then, unplug the interface board assembly.



3. Unplug the cables on the interface board. Then unscrew the two M3×6 screws and remove the USB_Hub board.



-
4. Tear the insulating pad off the interface board. Unscrew the two M3×6 screws on the DVI interface board. Then unscrew the two screws beside the DVI socket, the two screws beside micro-D socket and the two M2.5×6 screws.



NOTE

-
- Be careful not to damage the insulation between the DVI interface board and the USB_Hub board. If it is damaged, stick a new one.
-

5.4 Removing the SMR Assembly

1. First remove the 4 screw covers and then unscrew the 4 M3×8 screws.



2. Pull the left- and right-side boards outwards. Be sure to place the rubber ring in position when reassembling the right-side board.



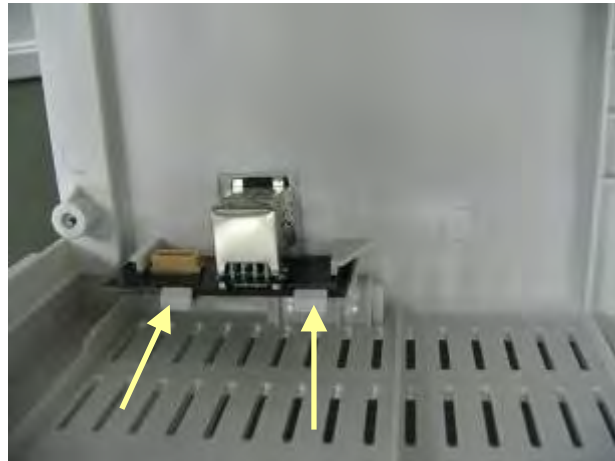
-
3. From the left side, remove the cable that connects the SMR interface board and the SMR communication board. Then take out the SMR from its housing.



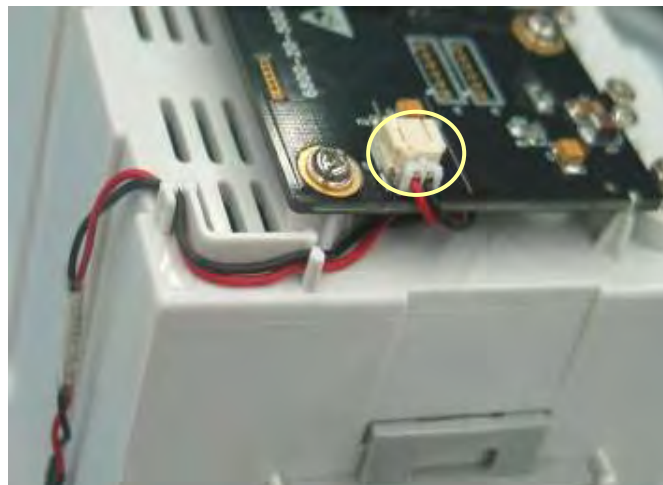
SMR

Housing

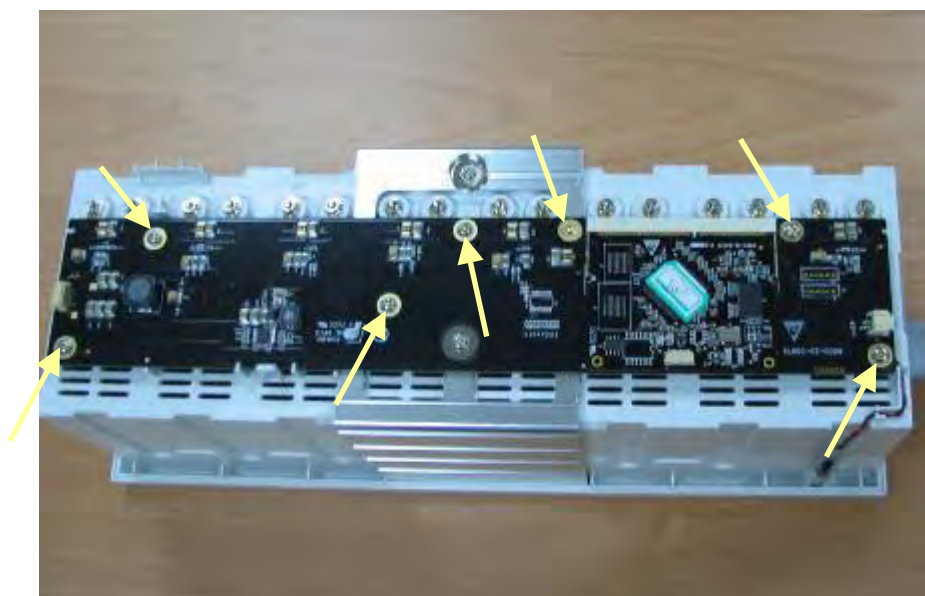
4. Release the two clips and take out the SMR interface board. Be sure not to damage the snap slot on the left side.



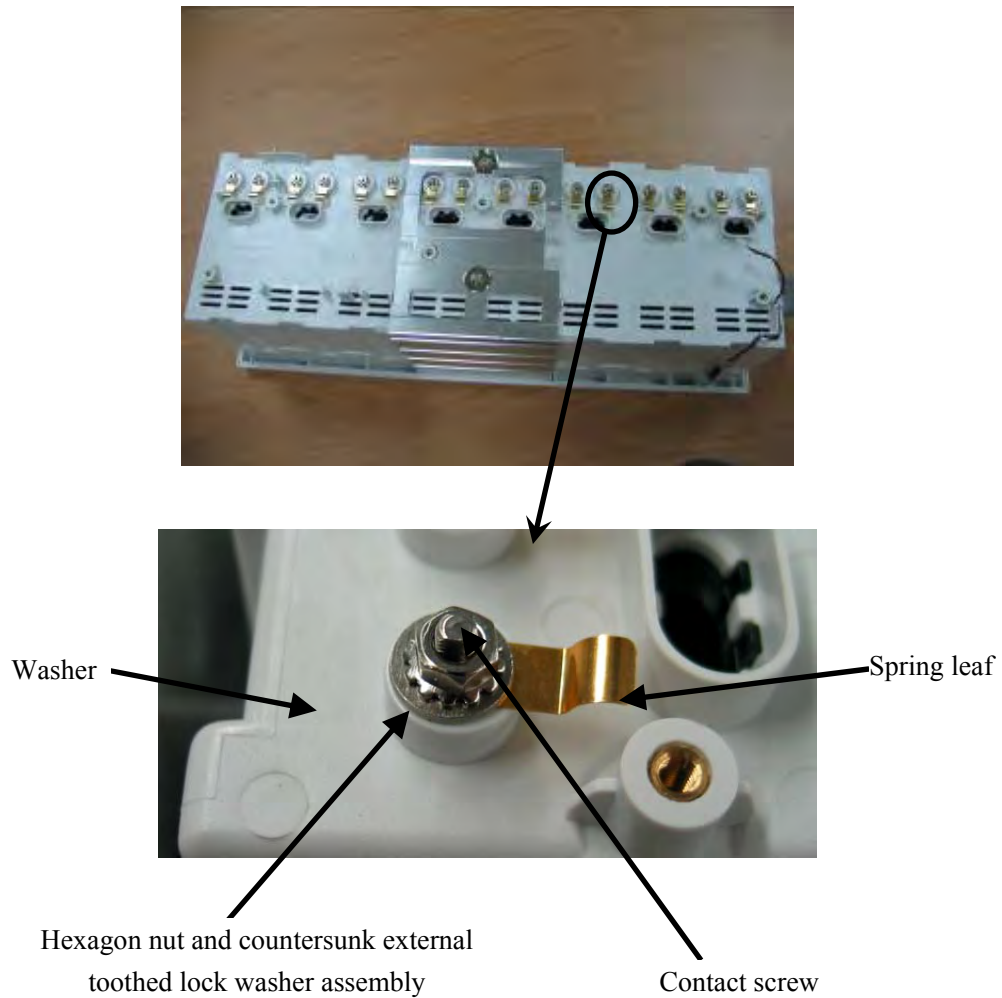
5. Remove the LED indicator, the light conductor and the cable that connects the SMR communication board and the LED board.



-
6. Release the clips and take out the Nios II module. Then unscrew the six M3×6 screws and remove the SMR communication board.



-
7. Use a socket wrench to unscrew the hexagon nut and countersunk external toothed lock washer assembly which can be further separated into the washer, springlet and contact screw.



5.5 Disassembling Modules

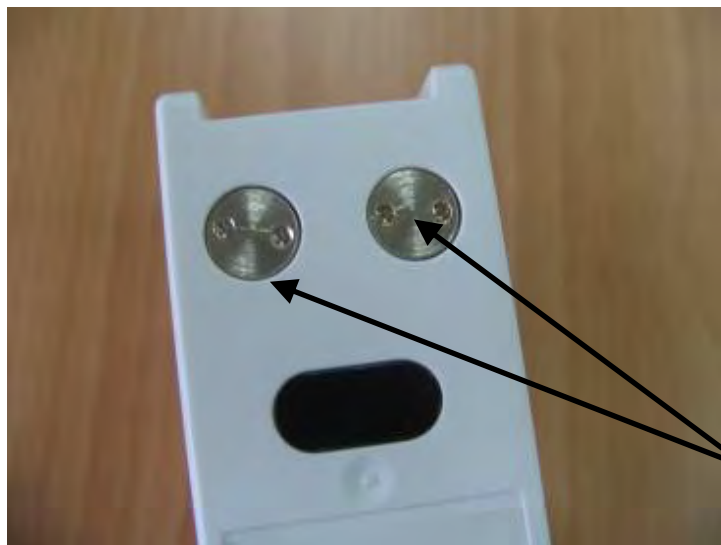
WARNING

- **For a reassembled module, a patient leakage current test must be performed before it is used again for patient monitoring.**
 - **Make sure the monitor is off before disassembling the modules.**
-

The monitor supports three types of modules, single-slot modules, 2-slot modules and 3-slot modules. The following part describes two typical disassembling procedures.

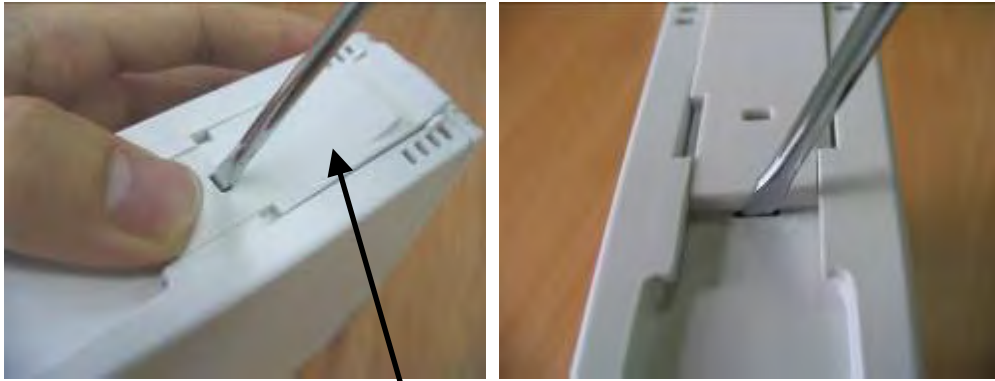
5.5.1 Disassembling the ICG Module

1. Unscrew the two contact screws and $\phi 3$ spring washers on the back of the module with a sharp nose pliers.



Contact screws

-
2. Unlock the snap close by pressing it down about 1 mm with a flat screwdriver. At same time, push the snap plate forward with your thumb until the snap close separate from its counterpart. Lift the snap plate with the flat screwdriver and remove it from the ICG module. In case the snap close relocks the snap plate when the end of the snap plate reaches the snap close, push the snap plate forward softly with the flat screwdriver.

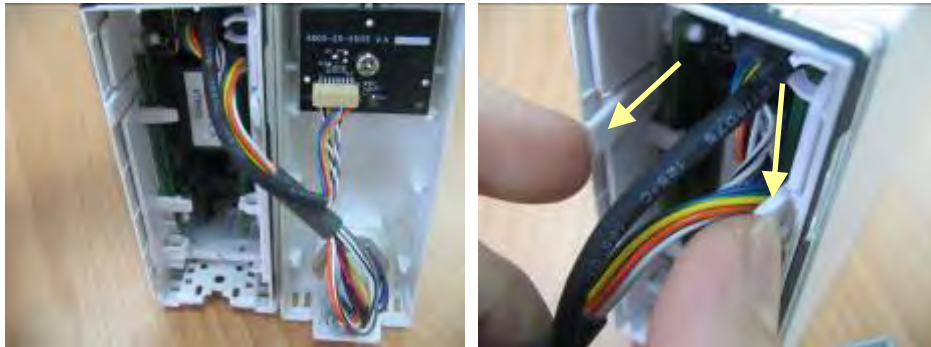


Snap plate

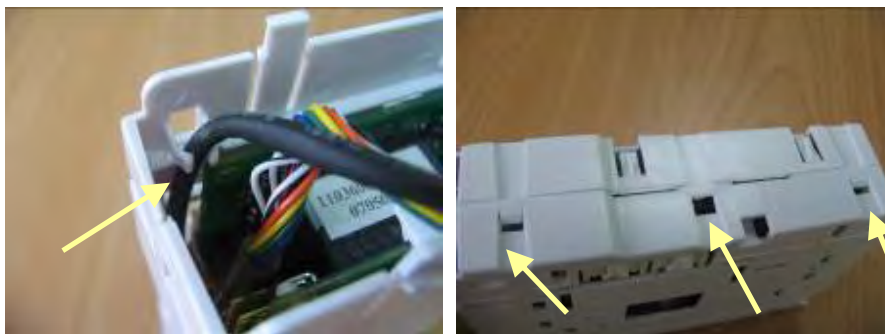
3. Unscrew the M3×6 screw. Then, press down, in turn, the two clips that engage the front panel. At same time, separate the front panel from the module's outer housing.



-
4. Remove the front panel carefully to avoid damaging the cables. The ICG module has an inner housing and an outer housing. Press the two clips on the inner housing about 1mm inwards to separate the inner and outer housing. Then, take off the outer housing.



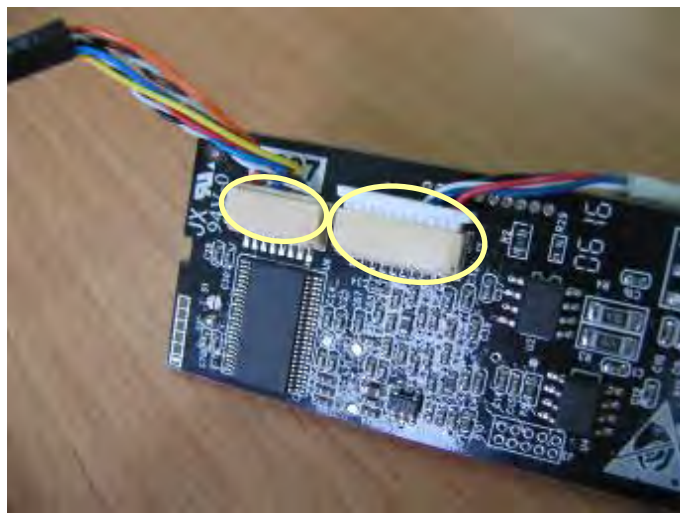
5. The inner housing consists of a left and a right side cover. Release the three clips as shown in the figure below to separate the two side covers.



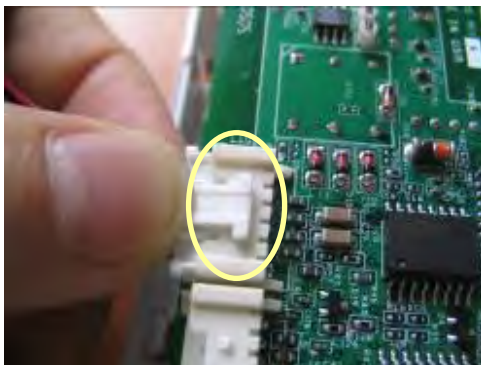
-
6. Take out the non-isolated power board and disconnect all the cables from it.



7. Take out the infrared communication board and disconnect all the cables from it.



8. Disconnect all the cables from the PCBA and remove the front panel with cables on it.



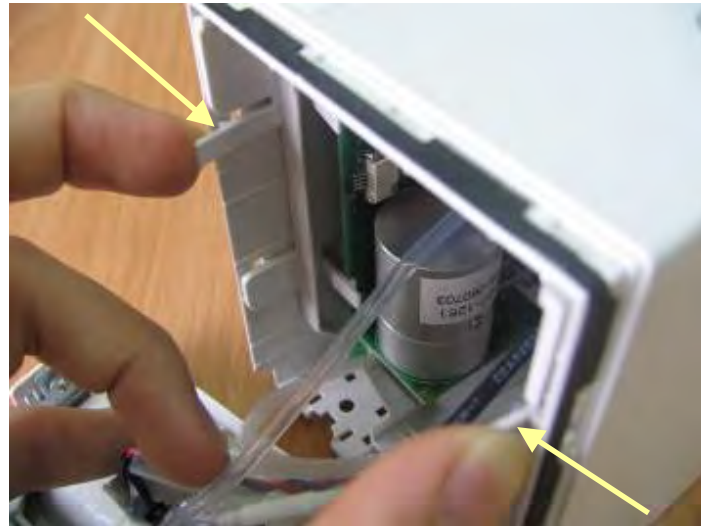
5.5.2 Disassembling CO₂ Module

1. Remove the four contact screws on the back and the two snap plates at the bottom and then unscrew the two M3×6 screws by referring to the procedure as described in **5.5.1 Disassembling the ICG Module**.

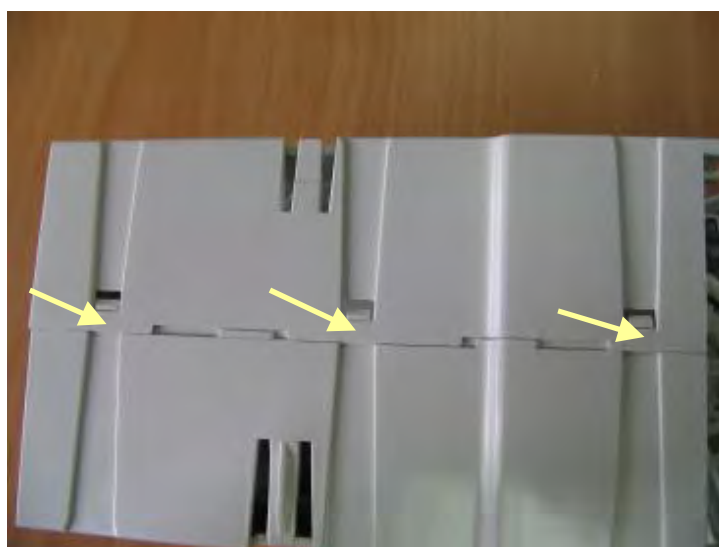


2. Remove the module's front panel by referring to the procedure as described in **5.5.1 Disassembling the ICG Module**. Then, press the two clips on the inner housing about 1mm inwards to separate the inner and outer housing. Then, take off the outer housing.





3. Hold the inner housing with both hands. Press the snap side with your thumb and push the other side with the index finger and the middle finger to separate the inner housing into two halves.

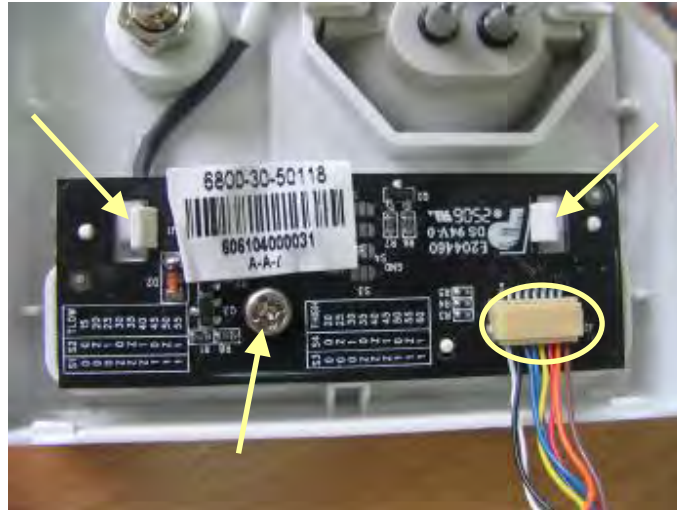




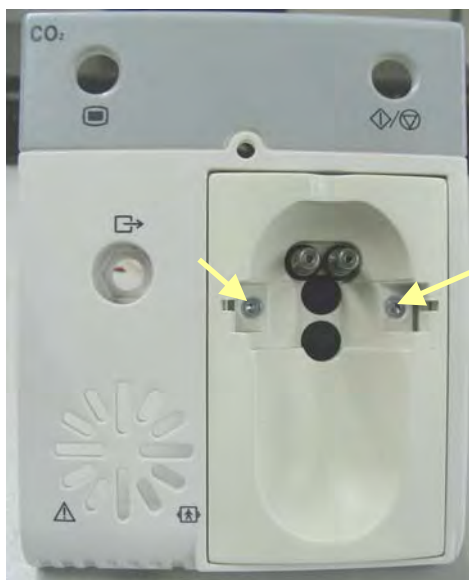
4. Remove the module infrared communication backboard and disconnect all the cables from it.



-
5. Disconnect the cables from the button board and unscrew the PT3×8 screw. Then release the two clips and take out the button board. After that, disconnect the cable to the fan and remove the button board.



-
6. Unscrew the two screws on the front panel. Disconnect the tubing between the watertrap assembly and CO₂ parameter board. Disconnect the cable between the watertrap and CO₂ parameter board. Then take out the watertrap connector assembly.



-
7. Disconnect the tubing between the CO₂ parameter board and the front panel. Then release the four clips and remove the CO₂ parameter board.

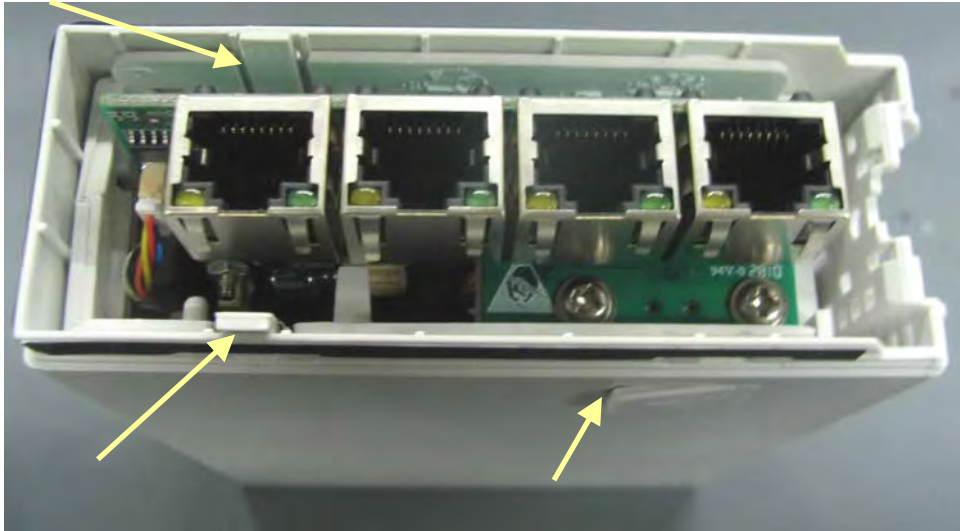


NOTE

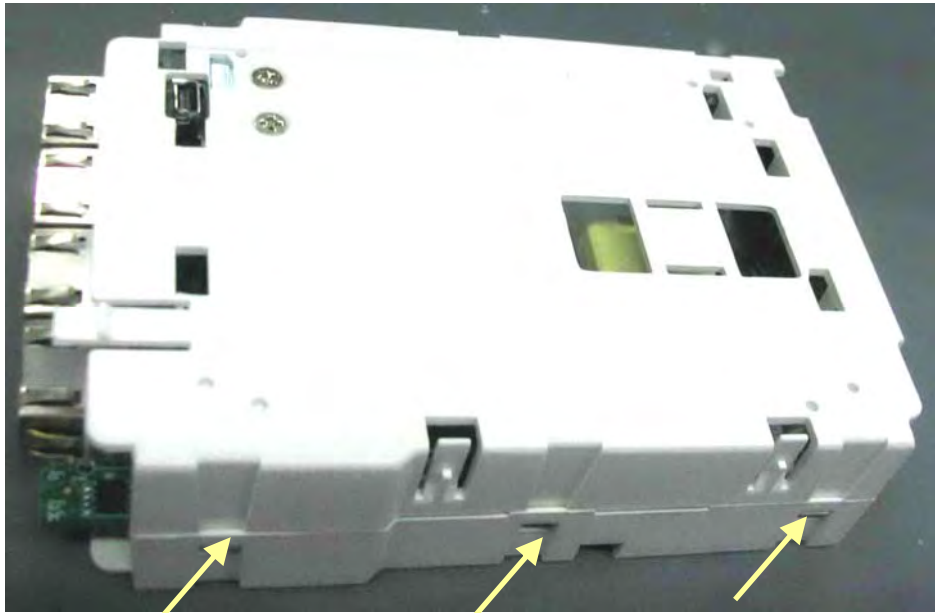
- When reinstalling, do not fold the tubings.

5.5.3 Disassembling the BeneLink Module

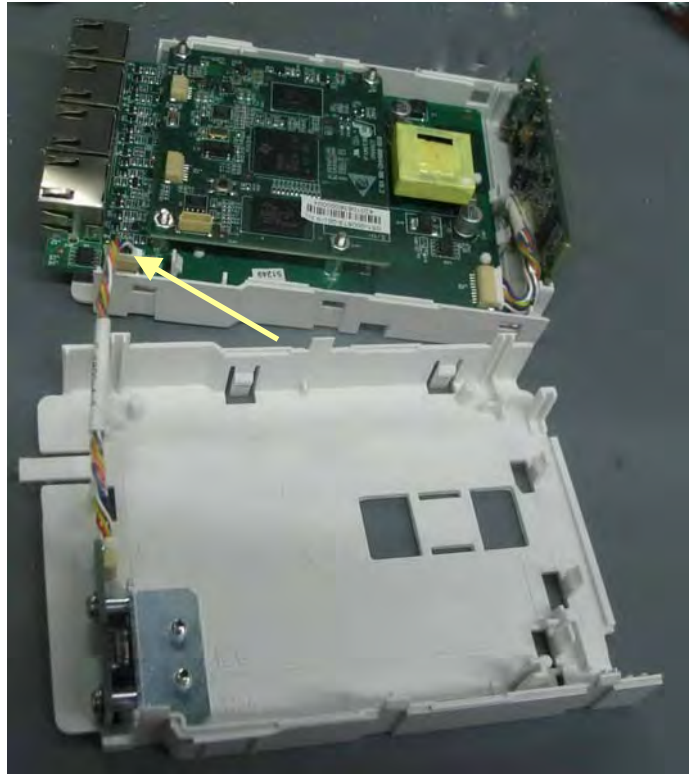
1. Remove the contact screws, the spanner, and the front cover by referring to steps 1 to 3 as described in *5.5.1 Disassembling the ICG Module*.
2. Take off the small cover board on one side of the rear cover. Then press the two clips about 1mm and take off the housing.



3. Release the three snaps to separate the two halves of the module side cover.



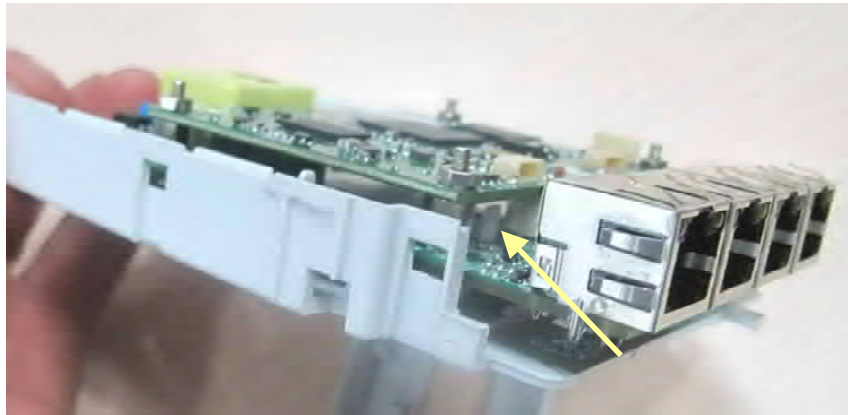
-
4. Take off the cable between the USB board and the interface board.



5. Take off the cable between the infrared communication board and the interface board to remove the infrared board.

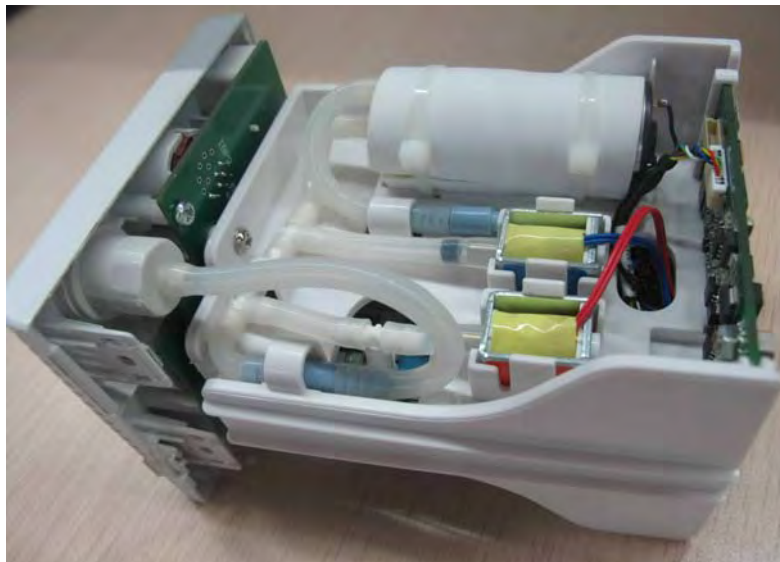
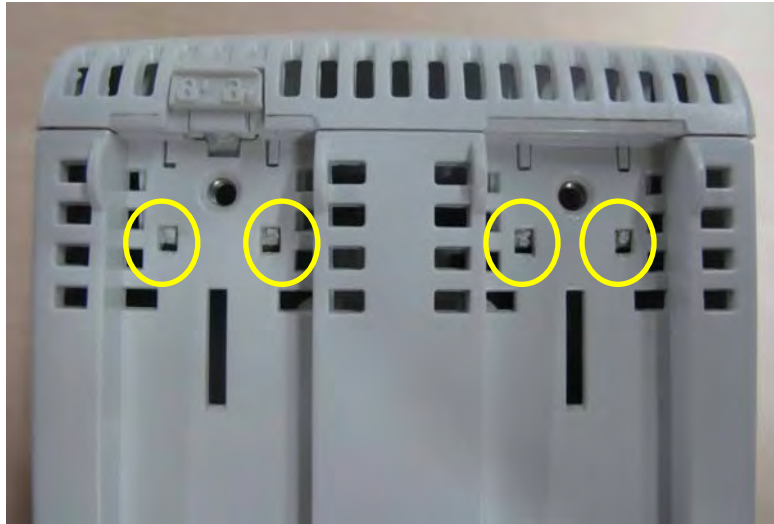


-
6. Unclench the four clips with a tweezer to remove the interface board.



5.5.4 Disassembling the New MPM Module

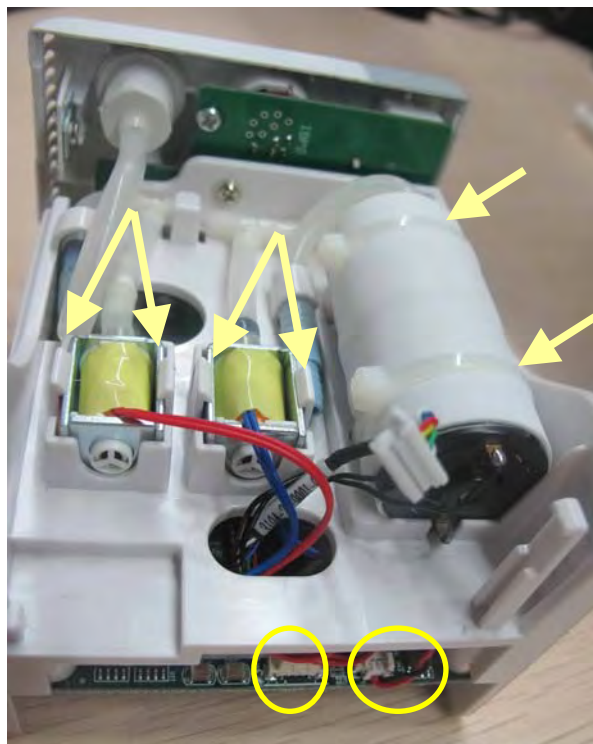
1. Remove the contact screws and the spanner by referring to steps 1 to 3 as described in *5.5.1 Disassembling the ICG Module*.
2. Unscrew the two M3×6 screws. Then press down, in turn, the four clips that engage the front cover with a small flat-bladed screwdriver. At the same time, pull off the rear cover.



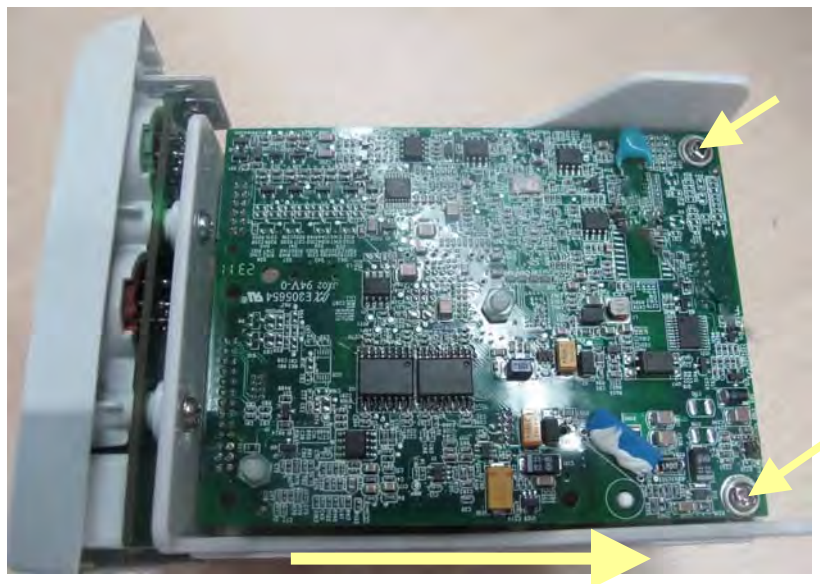
-
3. Disconnect the cable of the infrared communication board, and release the snap lock to remove the infrared communication board.



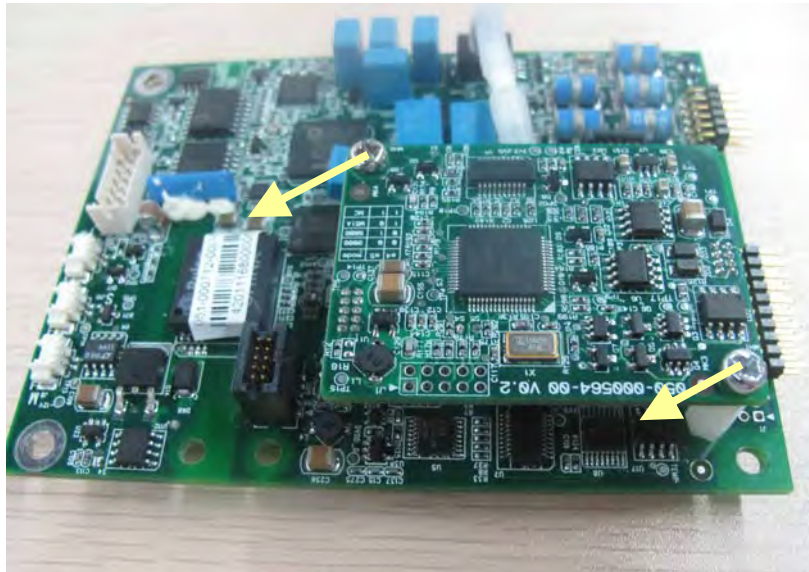
4. Disconnect the cables of the NIBP inflation pump and the relief valves from the parameter board. Snip off the cable ties to remove the pump. Release the snap locks to remove the valves.



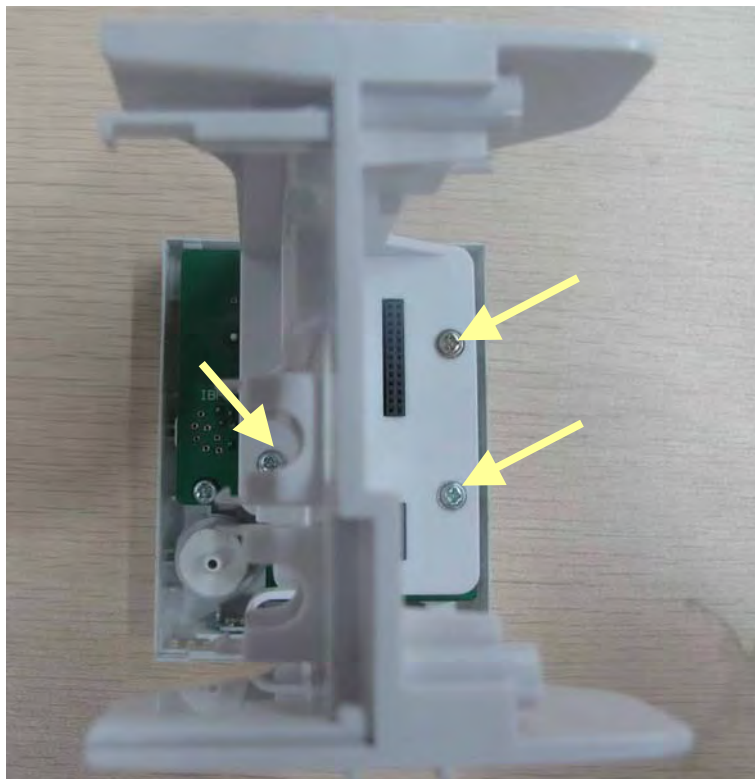
-
5. Unscrew the two M3×8 screws on the parameter board. Then pull off the parameter board rightward as shown below.



-
6. Unscrew the two M3×4 screws on the SpO₂ board to separate the SpO₂ board and the parameter board.



7. Unscrew the three M3×8 screws to separate the front panel assembly and the holder.



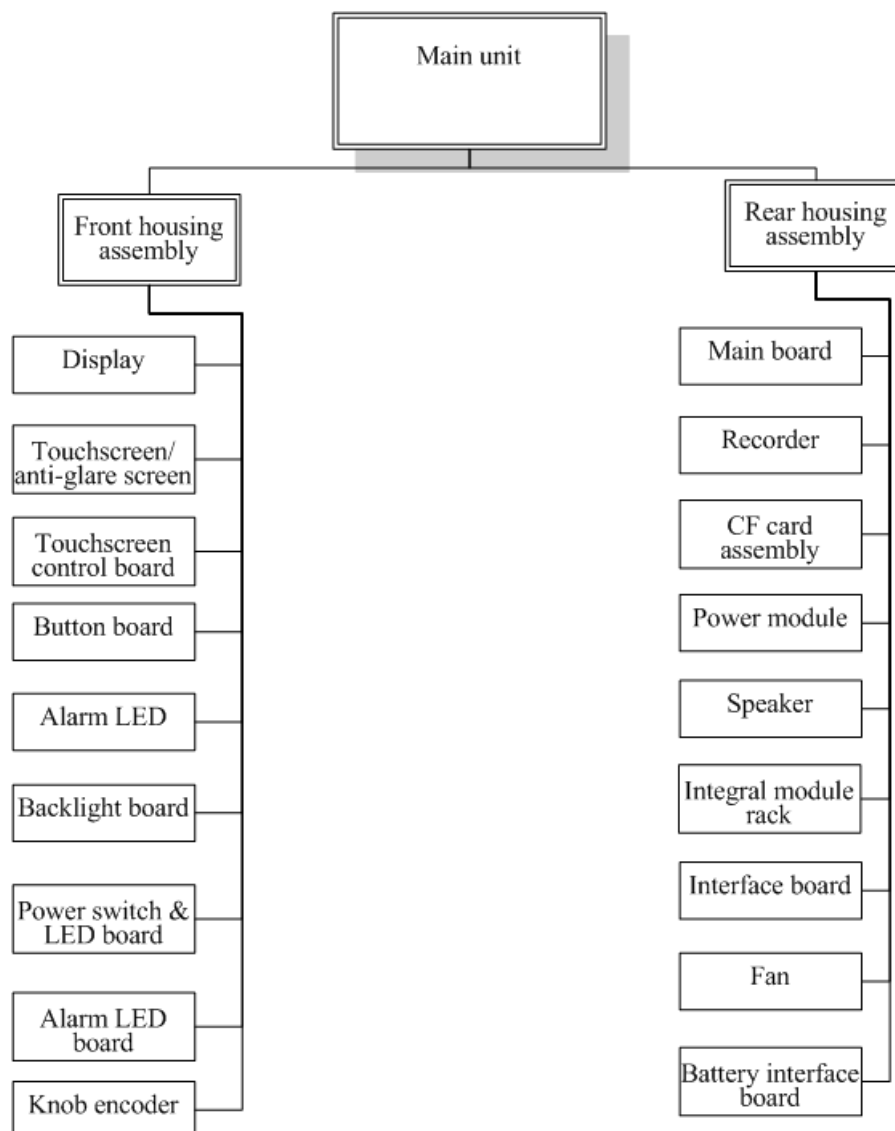
FOR YOUR NOTES

6 Parts

6.1 Introduction

This chapter contains the exploded views and parts lists of the main unit, satellite module rack and the parameter modules of the patient monitor. It helps the engineers to identify the parts during disassembling the patient monitor and replacing the parts.

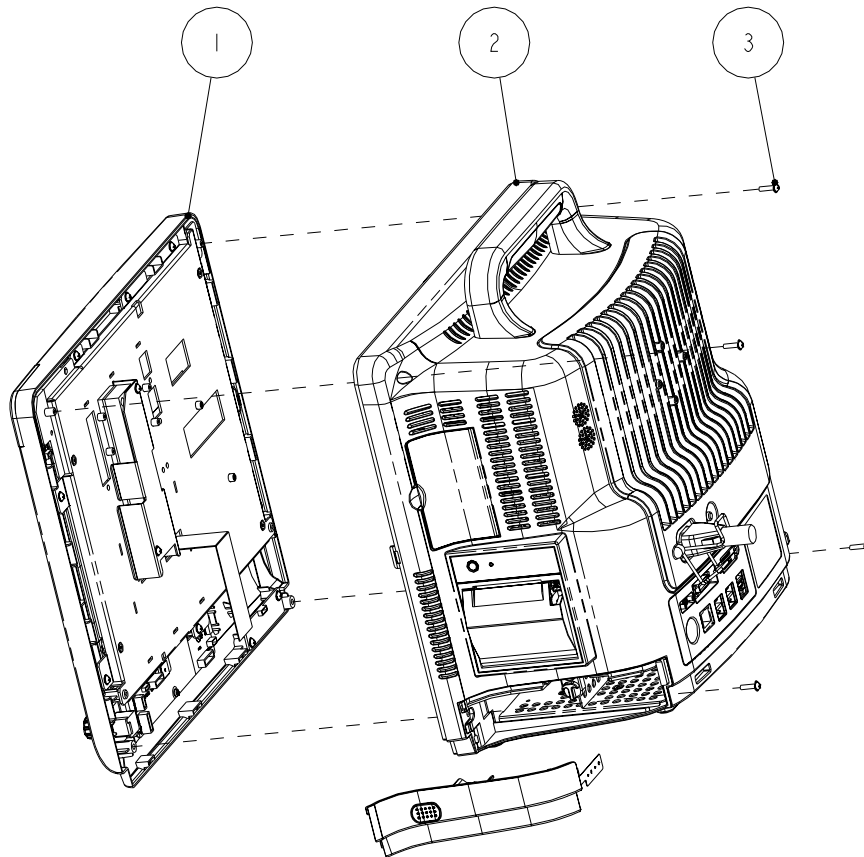
Hardware architecture of the main unit is shown below:



Remark: In the following tables, the P/N with the mark * is the specified P/N for DPM6 Standard configuration model.

6.2 Main Unit

6.2.1 Exploded View



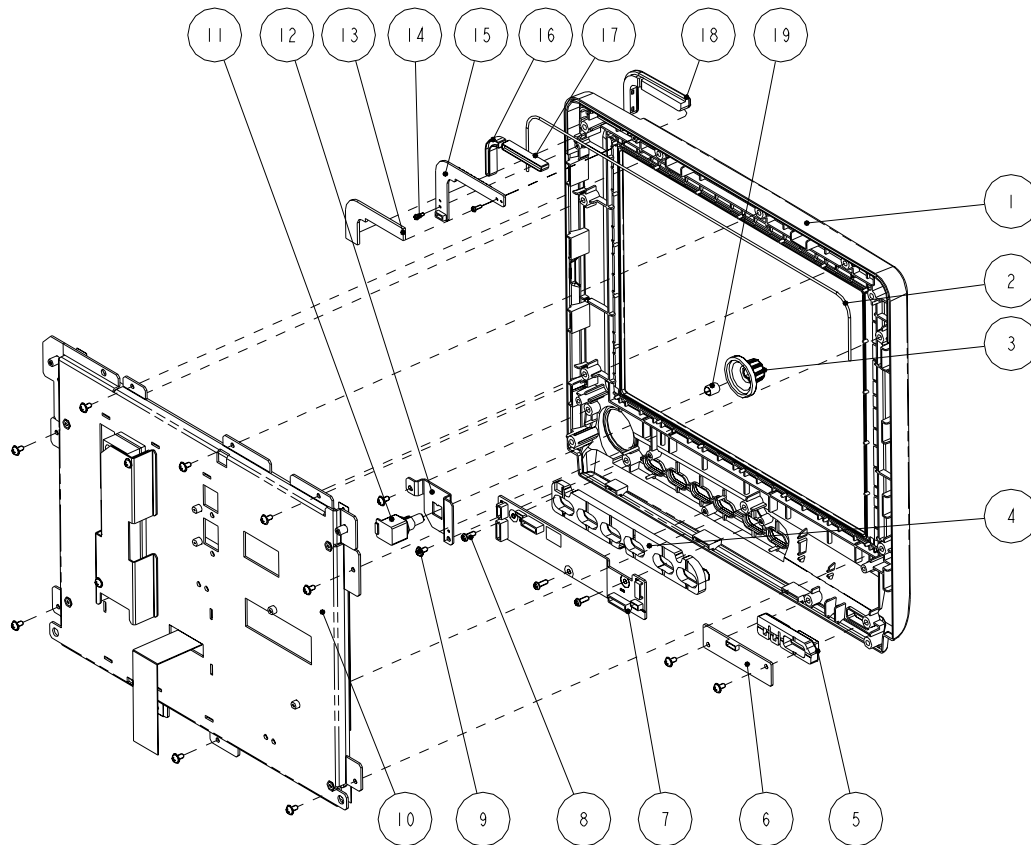
6.2.2 Parts List

SN	P/N	Description	Qty
1	6802-30-66761 or 6802-30-66762	Front housing assembly (with 12.1" LCD, anti-glare screen) Front housing assembly (with 12.1" LCD, touchscreen)	1
	*6812-30-66997 or *6812-30-66998	Front housing assembly (12.1" anti-glare screen) Front housing assembly (12.1" anti-glare screen)	
2	6802-30-66760 or *6812-30-66996	Main unit	1
3	M04-004017---	Crosshead screw M3 × 12	4

6.3 Front housing Assembly

6.3.1 12.1" LCD with Anti-glare Screen

Exploded View



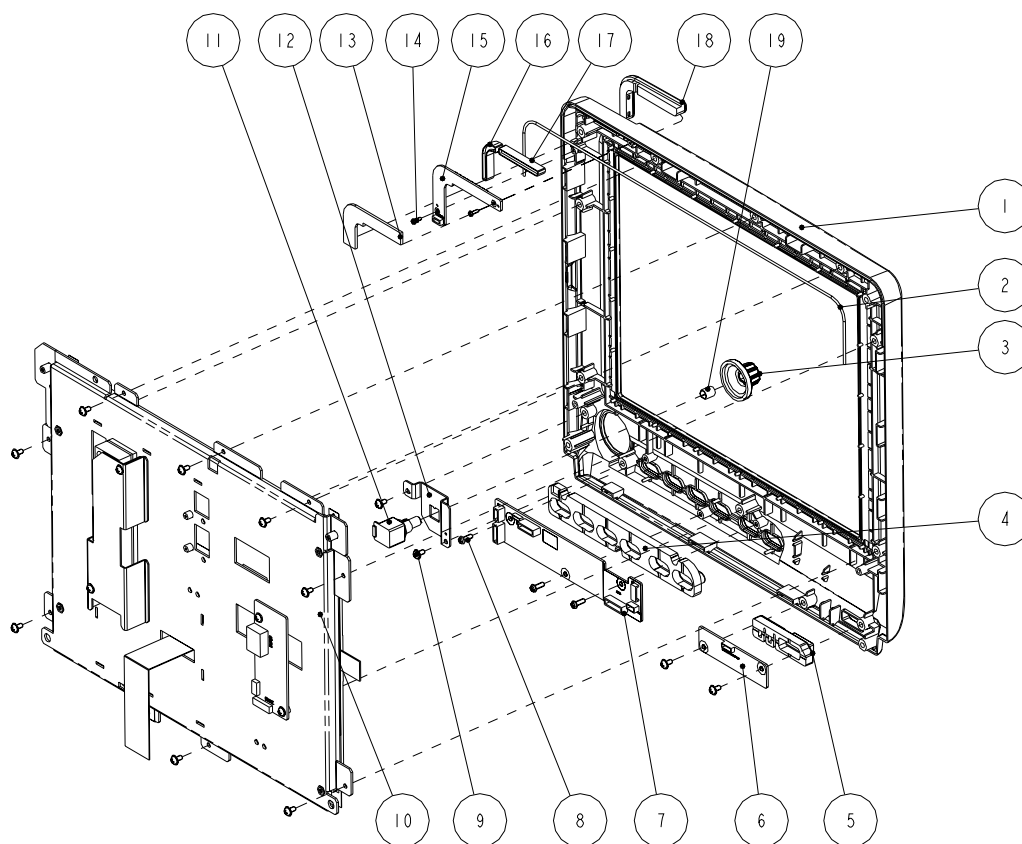
Parts List

SN	P/N	Description	Qty
1	043-000487-00 or 043-000487-01 or 043-000487-02 or *043-000490-00 or 043-000490-02	Front bezel (DPM6) (mold MR66687) Front bezel (DPM6/basic)	1
2	M6G-020015---	Hose	1
3	0651-20-76735	Knob encoder (mold MR76734)	1
4	6800-20-50198	Key pad	1
5	6802-20-66691-51	Power button (DPM6)	1
6	6802-30-66680	Power switch & LED board	1

7	6802-30-66812	Button board	1
8	M04-003105---	Tapping screw PT3×8	1
9	M04-004012---	Crosshead screw M3×6	13
10	6802-30-66901 or 6802-30-66772(optional)	12.1" LCD assembly (anti-glare screen)	1
11	0000-10-10789	Optical Encoder 16 steps 5VDC Dip6	1
12	042-000693-00	Encoder mounting board	1
13	6802-20-66729	Heat conductor for the LED cover	1
14	M04-051003---	Crosshead tapping screw PT2×6	1
15	6802-30-66682	LED board	2
16	6802-20-66690	Light conduction block 2	1
17	6802-20-66689	Light conduction block 1	1
18	6802-20-66688	Alarm LED cover	1
19	M6T-040001---	Clamp spring.1707P	1

6.3.2 12.1" LCD with Touchscreen

Exploded View

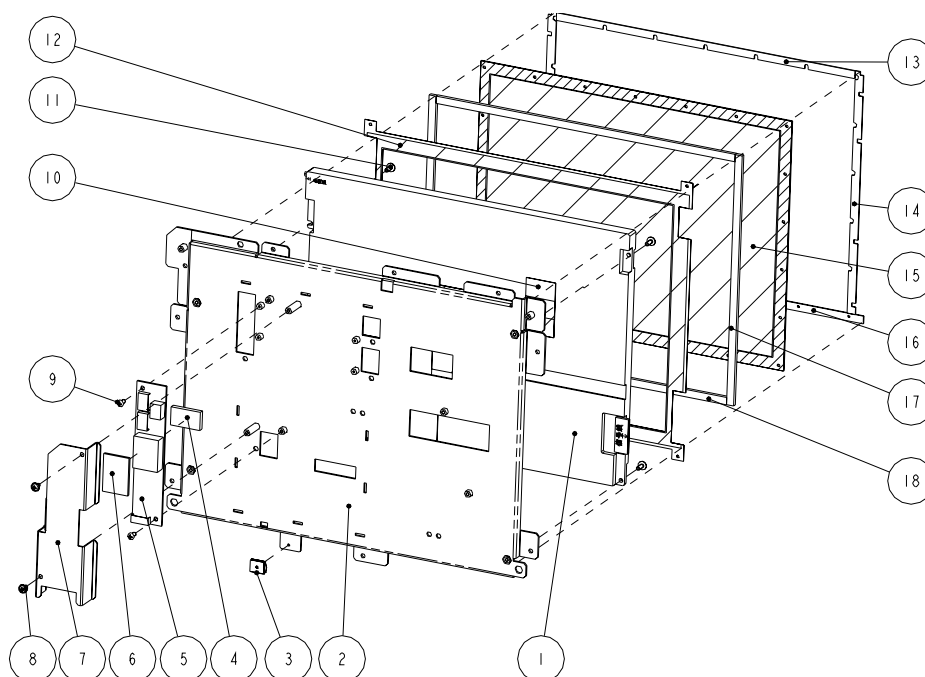


Parts List

SN	P/N	Description	Qty
1	043-000487-00 or 043-000487-01 or 043-000487-02 or * 043-000490-00 or 043-000490-02	Front bezel (DPM6) (mold MR66687) Front bezel (DPM6/basic)	1
2	M6G-020015---	Hose	1
3	0651-20-76735	Knob encoder (mold MR76734)	1
4	6800-20-50198	Key pad	1
5	6802-20-66691-51	Power button (DPM6)	1
6	6802-30-66680	Power switch & LED board	1
7	6802-30-66814	Button board (AU 12.1" screen G121SN01 + touchscreen)	1
8	M04-003105---	Tapping screw PT3×8	3
9	M04-004012---	Crosshead screw M3×6	13
10	6802-30-66902 or 6802-30-66773(optional)	12.1" LCD assembly (touchscreen)	1
11	0000-10-10789	Optical Encoder 16 steps 5VDC Dip6	1
12	042-000693-00	Encoder mounting board	1
13	6802-20-66729	Heat conductor for the LED cover	1
14	M04-051003---	Crosshead tapping screw PT2×6	2
15	6802-30-66682	LED board	1
16	6802-20-66690	Light conduction block 2	1
17	6802-20-66689	Light conduction block 1	1
18	6802-20-66688	Alarm LED cover	1
19	M6T-040001---	Clamp spring.1707P	1

6.3.3 12.1" Screen Assembly (with anti-glare screen)

Exploded View



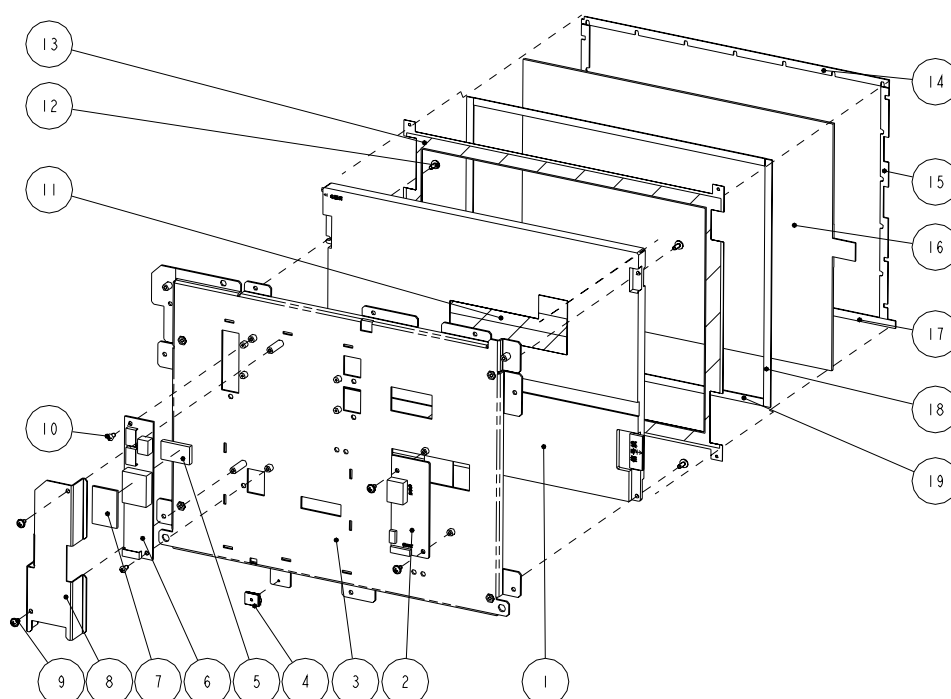
Parts List

SN	P/N	Description	Qty
1	0000-10-10752 or 0010-10-42633 (optional)	LCD TFT 12.1"	1
2	042-000784-00 or 042-000783-00	12" SHARP screen mounting board, or 12" AU screen mounting board	1
3	0000-10-10996	Beryllium copper leaf, 92-047, inferior smooth	1
4	042-000185-00	TPI inverter washer	1
5	022-000014-00 or 022-000001-00	TPI inverter	1
6	6800-20-50692	Backlight board heat-conductive insulator	1
7	042-000601-00	TPI inverter shield	1
8	M04-004012---	Crosshead screw with washer M3×6	2
9	M04-002505---	Crosshead screw GB/T818-2000 M3×6	2
10	6802-20-66731 or 6802-20-66650	12"SHARP screen cable or AU screen cable	1
11	M04-004015---	Crosshead screw with washer M3×8	4
12	6802-20-66795	Screen cover	1

13	6802-20-66694	Dust-proof strip 1	1
14	6802-20-66695	Dust-proof strip 2	2
15	6802-20-66693	Anti-glare screen	1
16	6802-20-66800	Anti-glare screen water-proof strip	1
17	6802-20-66741	Dust-proof strip 4	2
18	6802-20-66740	Dust-proof strip 3	2

6.3.4 12.1" Screen Assembly (with touchscreen)

Exploded View



Parts List

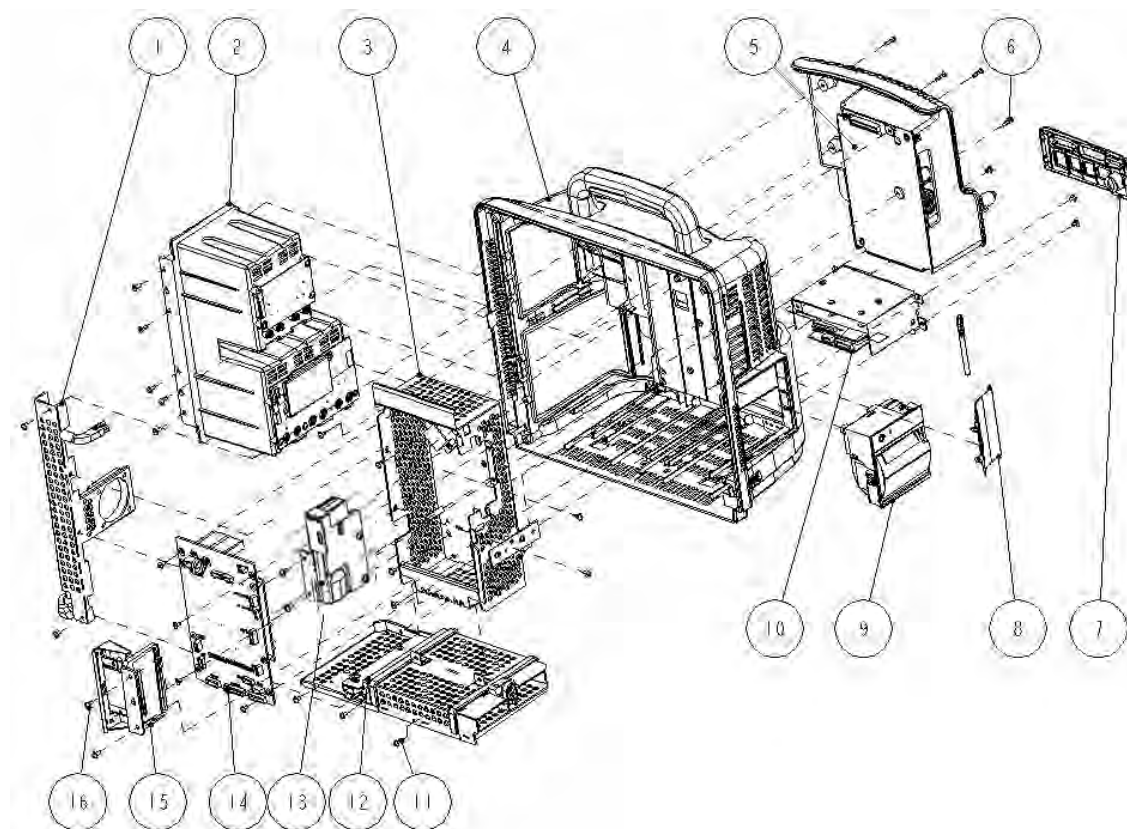
SN	P/N	Description	Qty
1	0000-10-10752 or 0010-10-42633 (optional)	LCD TFT 12.1"	1
2.	6800-30-50082	Touchscreen control board	1
3	042-000784-00 or 042-000783-00	12" SHARP screen mounting board, or 12" AU screen mounting board	1
4	0000-10-10996	Beryllium copper leaf, 92-047, inferior smooth	1
5	042-000185-00	TPI inverter washer	1
6	022-000014-00 or 022-000001-00	TPI inverter	1

7	6800-20-50692	Backlight board heat-conductive insulator	1
8	042-000601-00	TPI inverter shield	1
9	M04-004012---	Crosshead screw with washer M3×6	6
10	M04-002505---	Crosshead screw GB/T818-2000 M3×6	2
11	6802-20-66731 or 6802-20-66650	12"SHARP screen cable or AU screen cable	1
12	M04-004015---	Crosshead screw with washer M3×8	4
13	6802-20-66795	Screen cover	1
14	6802-20-66696	Touchscreen strip 1	1
15	6802-20-66697	Touchscreen strip 2	2
16	0000-10-10799	Touchscreen, 12.1"	1
17	6802-20-66801	Touchscreen water-proof strip	1
18	6802-20-66739	Touchscreen strip 4	2
19	6802-20-66738	Touchscreen strip 3	2

6.4 Main Unit

6.4.1 Main Unit Assembly

Exploded View

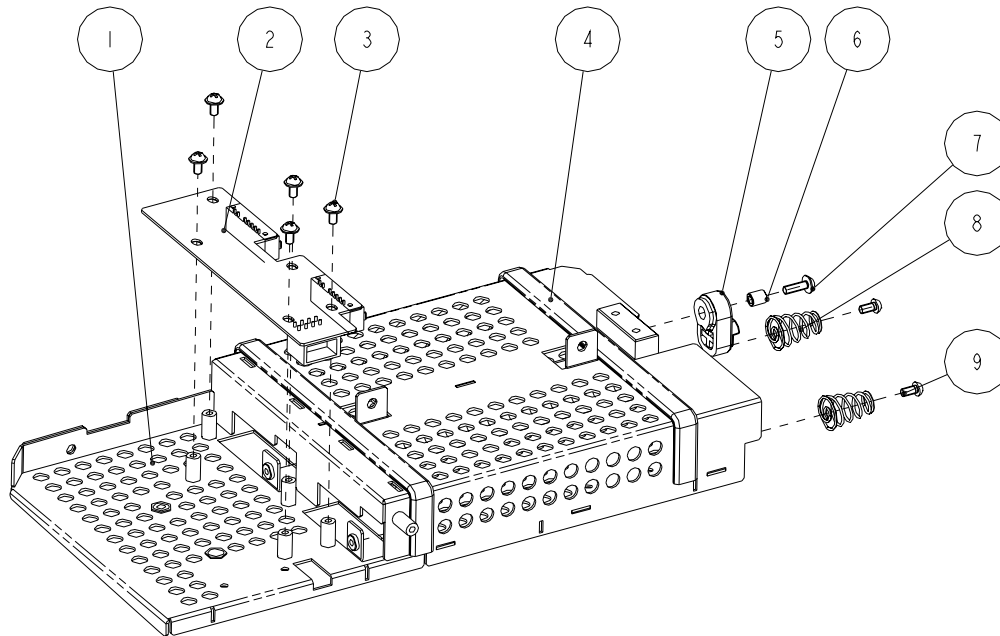


Parts List

SN	P/N	Description	Qty
1	6802-30-66768	Fan assembly	1
2	6802-30-66765	Integral module rack	1
3	115-003229-00	Main support assembly	1
4	6802-30-66766	Rear housing assembly	1
5	6802-30-66770	Power module assembly	1
6	M04-051014---	Crosshead screw M3×12	4
7	043-000059-00 or 043-000059-01 or *6812-20-67000-51 or 043-004115-00 or 043-000472-00	Interface board cover(new interface) Interface board cover (basic) Interface board cover	1
8	6802-20-66707-51 or 043-004116-00	CF card door	1
9	TR6F-30-67306 or 115-001290-00	TR6F recorder	1
10	6802-30-66769 or *6812-30-66994 or 115-003230-00	Interface board assembly	1
11	M04-004012---	Crosshead screw M3×6	22
12	6802-30-66763	Battery compartment assembly	1
13	115-003228-00 115-015189-00	6802 internal wireless AP assembly 6802 new internal wireless AP assembly (ASUS)	1
14	6802-30-66767 or *6812-30-66993 or 115-003231-00	Master control board assembly	1
15	115-001906-00 or *6812-30-66995	6802 CF card assembly (211 driving board) CF card slot cover kit	1
16	M04-004015---	Crosshead screw M3×8	7

6.4.2 Battery Compartment Assembly

Exploded View

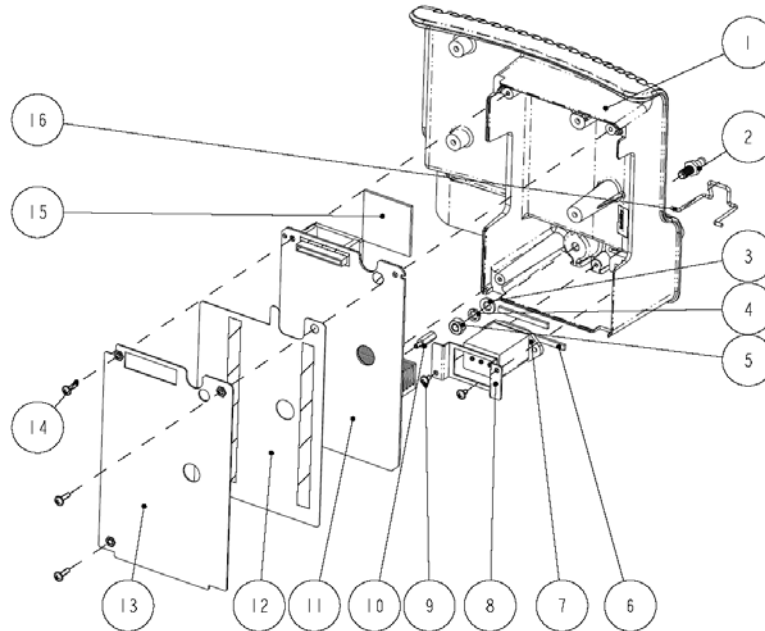


Parts List

SN	P/N	Description	Qty
1	6802-20-66714	Battery compartment	1
2	6802-30-66653	Battery interface board	1
3	M04-004012---	Crosshead screw M3×6	5
4	M6T-030008---	Conducting block 9.5×3.2	2
5	6802-20-66715 or 043-004095-00	Battery latch	1
6	6802-20-66798	Battery latch fixture	1
7	M04-004013---	Crosshead screw M3×10	1
8	DA8K-20-27052	Spring	2
9	M04-002505---	Crosshead screw M3×6	2

6.4.3 Power Module assembly

Exploded View

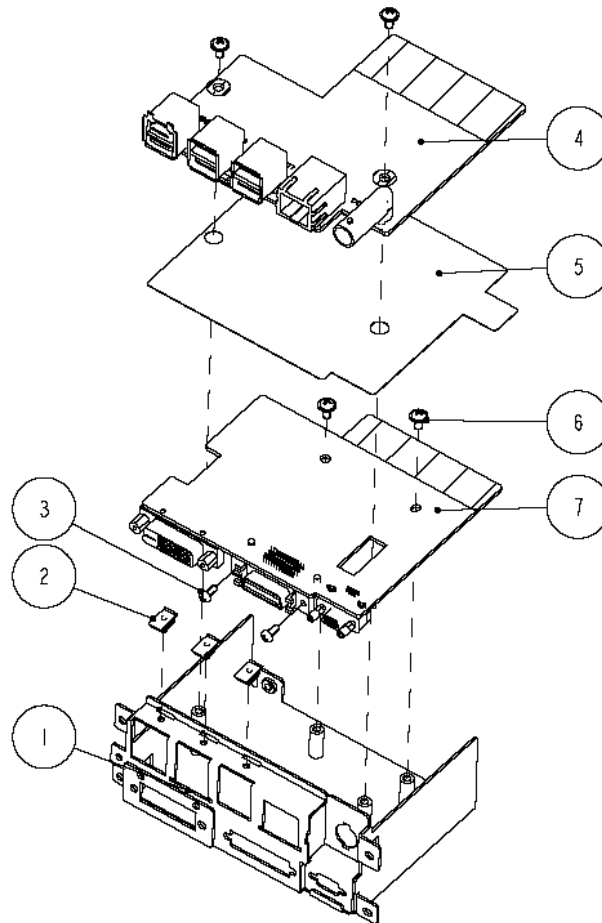


Parts List

SN	P/N	Description	Qty
1	6802-20-66719-51 or 044-000446-00	Power supply compartment(mould MR66719)	1
2	0509-20-00098	Grounding pole	1
3	M04-021003---	Flat washer, GB97.1 6	1
4	M04-004504---	Spring washer, GB93 6	1
5	M04-004401---	Stainless steel nut, GB6170 M6	1
6	6800-20-50636	Waterproof strip for power socket	2
7	009-000255-00	AC input socket and cable	1
8	6800-20-50218	power socket fixture	1
9	M04-004012---	Crosshead screw M3 × 6	2
10	M90-000136---	Plastic double-ended bolt M3 × 15+6-8	1
11	6802-30-66651	Power board	1
12	6802-20-66721	Insulating plate	1
13	6802-20-66720	Power supply cover	1
14	M04-004017---	Crosshead screw M3 × 12	3
15	6802-20-66796	Power board heat conductor	1
16	9211-20-87369	Pothead for power supply plugs	1

6.4.4 Interface Board Assembly

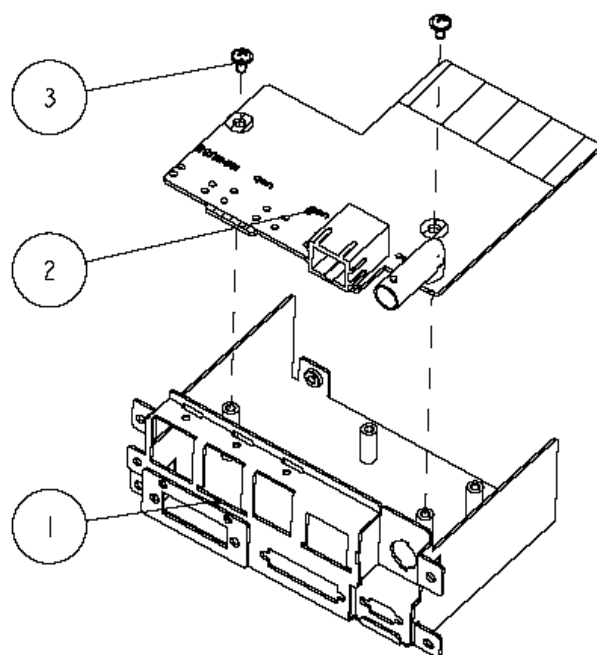
6.4.4.1 Interface Board Assembly (6802-30-66769)



Parts List

SN	P/N	Description	Qty
1	042-000059-00	Interface board frame (new interface)	1
2	M6T-030006---	Leaf	3
3	M04-051045---	Crosshead screw M2.5×6	2
4	051-000020-00	USB_Hub board (no analog output)	1
5	047-000128-00	Interface board insulating plate	1
6	M04-004012---	Crosshead screw M3×6	4
7	051-000019-00	DVI interface board (can support IABP)	1

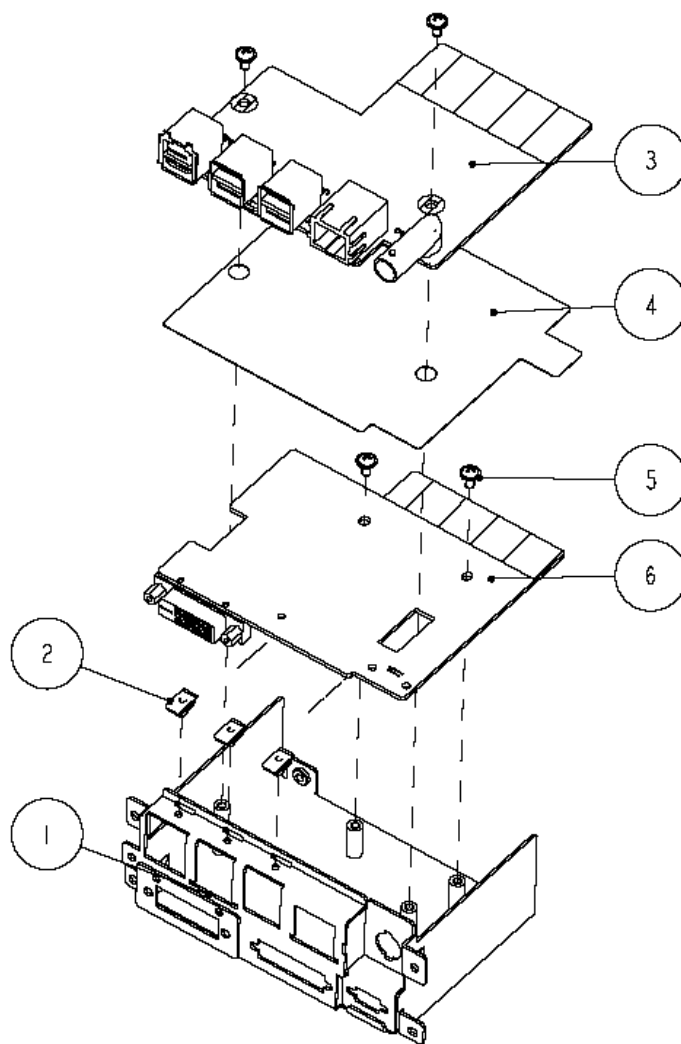
6.4.4.2 Interface Board Assembly (6812-30-66994)



Part List

SN	P/N	Description	Qty
1	042-000059-00	Interface board frame (new interface)	1
2	6812-30-66999	6812 network interface board	1
3	M04-004012---	Cross pan head screw with washer M3×6	2

6.4.4.3 Interface Board Assembly (115-003230-00)

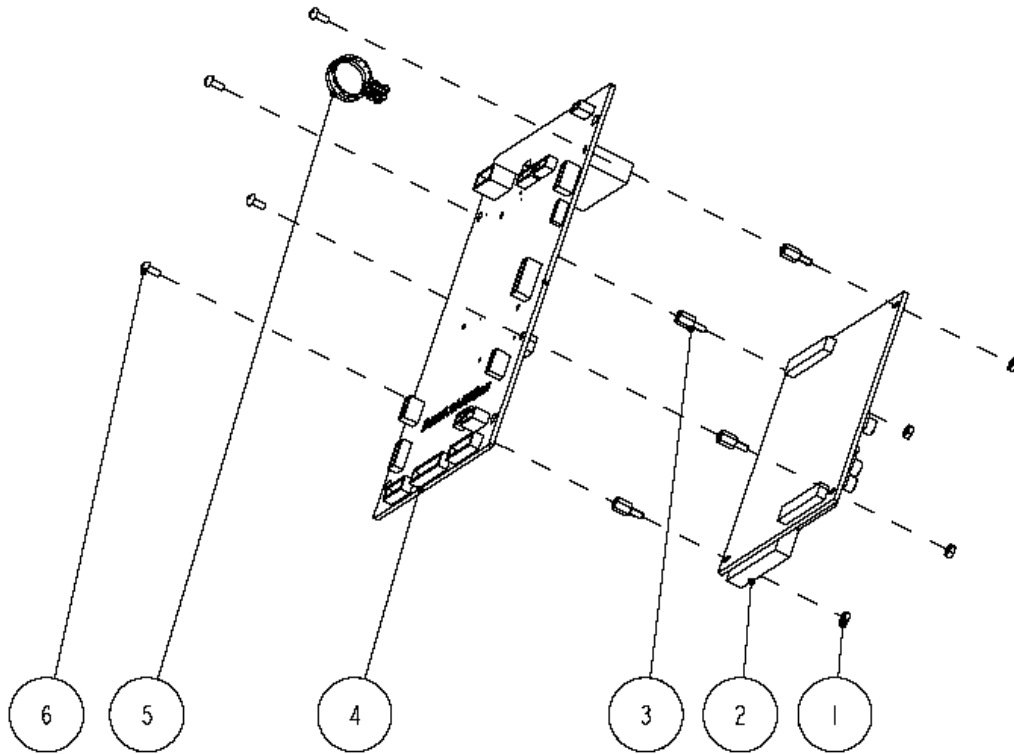


Part List

SN	P/N	Description	Qty
1	042-000059-00	Interface board frame (new interface)	1
2	M6T-030006---	Leaf	3
3	051-000020-00	USB_Hub board (no analog output)	1
4	047-000128-00	Interface board insulating plate	1
5	M04-004012---	Cross pan head screw with washer M3×6	4
6	051-000019-01	DVI interface board (Standard configuration)	1

6.4.5 Main Board Assembly

Exploded View

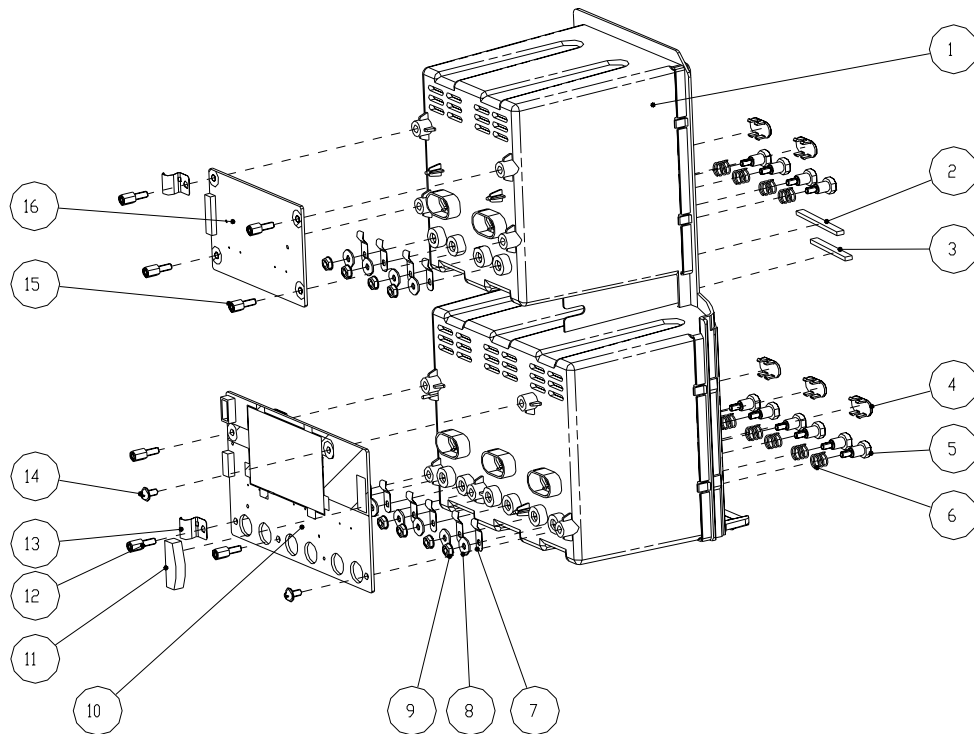


Parts List

SN	P/N	Description	Qty
1	M04-011001---	Nut, GB6170 M2.5	4
2	6800-30-50058 or *6812-30-66989	MCF547x CPU module (basic configuration/non-lead process) MCF547x CPU module (+standard identifying code)	1
3	M04-060021---	Screw M2.5x7+6-6	4
4	051-000223-00 or 051-000223-01	6802 mother board (full configuration) 6802 mother board (standard configuration)	1
5	6802-20-66724 or 043-004100-00	PCB handle	1
6	M04-051045---	Crosshead screw M2.5×6	4

6.4.6 Integral module rack

Exploded View



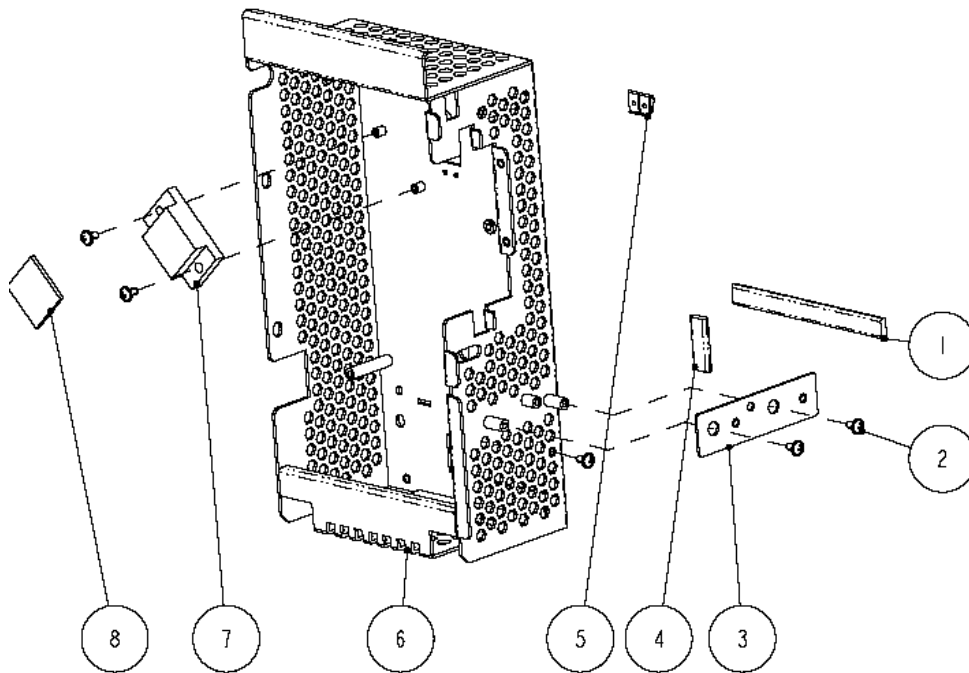
Parts List

SN	P/N	Description	Qty
1	6802-20-66709 or 043-004101-00	Integral module rack	1
2	6802-20-66817	Waterproof strip 2 for integral module rack	1
3	6802-20-66816	Waterproof strip 1 for integral module rack	1
4	6800-20-50279	Infrared light filter	5
5	6800-21-51100	Contact screw	10
6	6800-20-50261	Contact spring	10
7	6800-20-50388	Leaf	10
8	M04-021024---	Flat washer, GB96 3	10
9	M04-011002---	M3 nut with spring washer	10
10	051-000244-00	6802 3-slot communication board	1
11	9200-20-10512	Rubber foot	1
12	M04-060026---	Stud screw M3×7+6-6	5
13	6802-20-66710	Spring coil	2

14	M04-004012---	Pan head screw M3×6	2
15	M04-000206---	Stud screw M3×6+8-5	2
16	6800-30-50073	MPM module rack communication board	1

6.4.7 Main Support Assembly

Exploded View

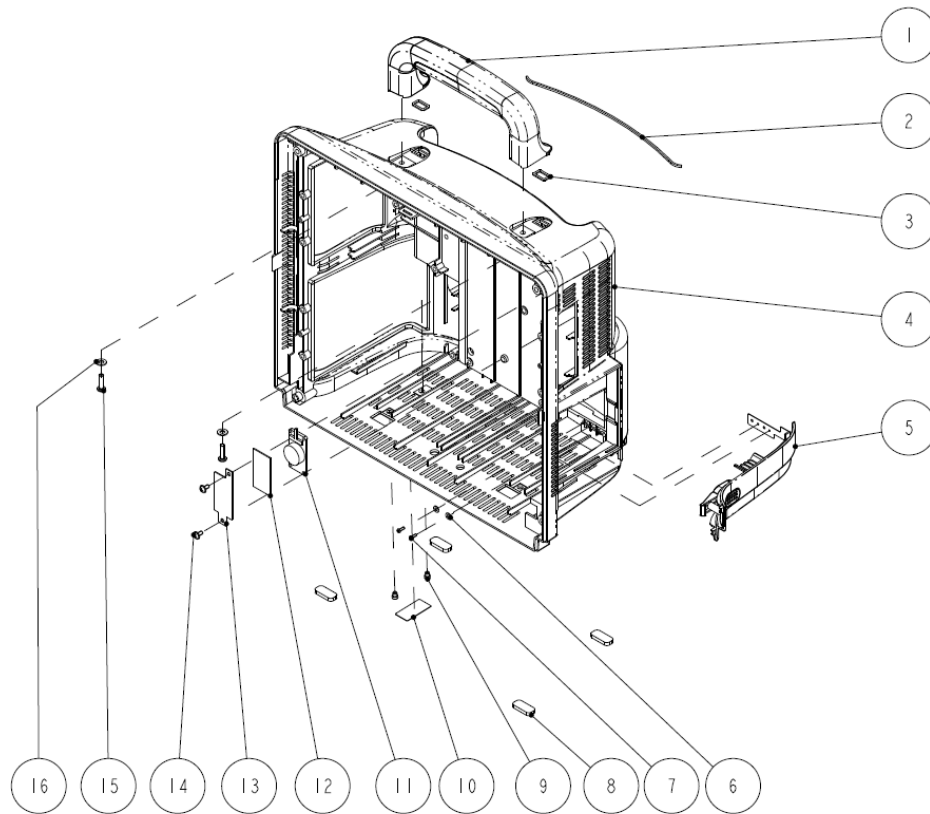


Parts List

SN	P/N	Description	Qty
1	M6T-030008---	Conducting block 9.5×3.2	1
2	M04-004012---	Crosshead screw M3×6	5
3	6802-20-66718	Recorder fixing plate	1
4	M6T-030008---	Conducting block 9.5×3.2	1
5	M6T-030006---	Leaf	2
6	6802-20-66717	Main support	1
7	6802-20-66789	CPU radiator	1
8	6802-20-66793	CPU heat conductor	1

6.4.8 Rear Housing Assembly

Exploded View



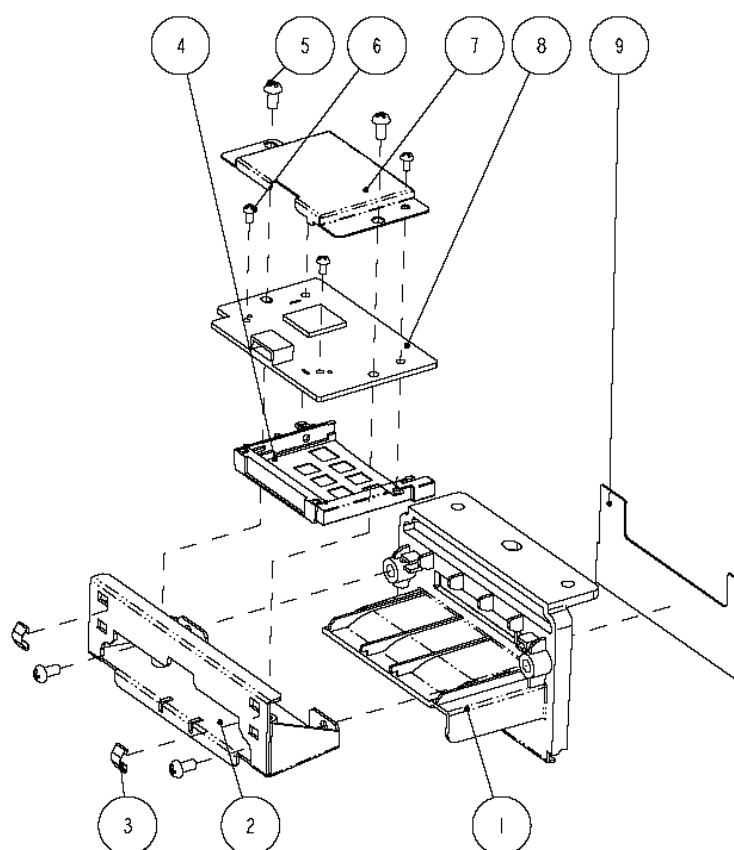
Parts List

SN	P/N	Description	Qty
1	6802-20-66699 or 043-004099-00	Handle	1
2	6802-20-66737	Waterproof strip for the power supply	1
3	6802-20-66736	Handle waterproof pad	2
4	6802-20-66698 or 043-004091-00	Rear housing	1
5	6802-30-66764	Battery door assembly	1
6	M04-021000---	Flat washer, GB97.1 2.5	2
7	M04-051003---	Crosshead tapping screw PT2×6	2
8	6800-20-50233--	Cushion	4
9	049-000518-00	6802 screw rubber	2
10	047-009145-00	Installing bolt lable (EN)	1

SN	P/N	Description	Qty
11	6802-20-66668	Speaker and cables	1
12	8002-20-36218	Crosshead screw M3×6	1
13	6802-20-66791	Speaker cover	1
14	M04-004012---	Cross pan head screw with washer M3×6	2
15	M04-051085---	Tapping screw PT4×14	2
16	M04-004702---	Flat washer, GB97.1 4	2

6.4.9 CF Card Assembly

CF Card Assembly (115-001906-00)

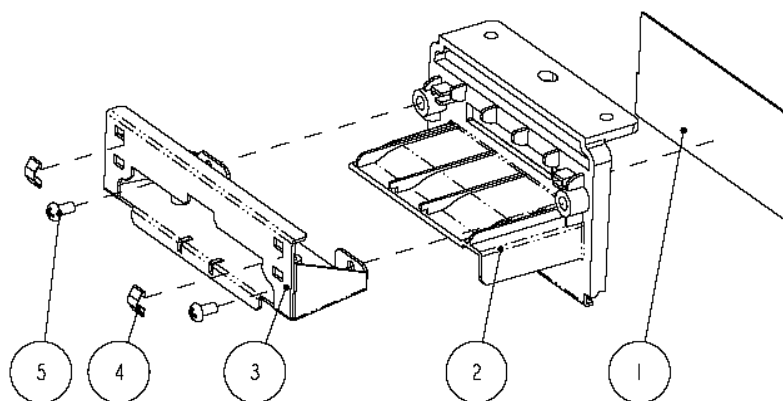


Parts List

SN	P/N	Description	Qty
1	6802-20-66725 or 043-004098-00	Card cover	1
2	6802-20-66726	CF card board fixture	1
3	M6T-030005---	Leaf, 178S30(1EA=406mm)	2

4	M39-000234---	Ejector for CF card socket (50pin)	1
5	M04-002505---	Crosshead screw M3×6	4
6	M04-051137---	Crosshead screw M2×4	4
7	042-000317-00	CF card shield	1
8	051-000104-00	9211 CF card board (shield can be installed)	1
9	6802-20-66674 or 047-000731-00	WLAN overlay Slot cover overlay	1

CF Card Slot Cover Kit (6812-30-66995)

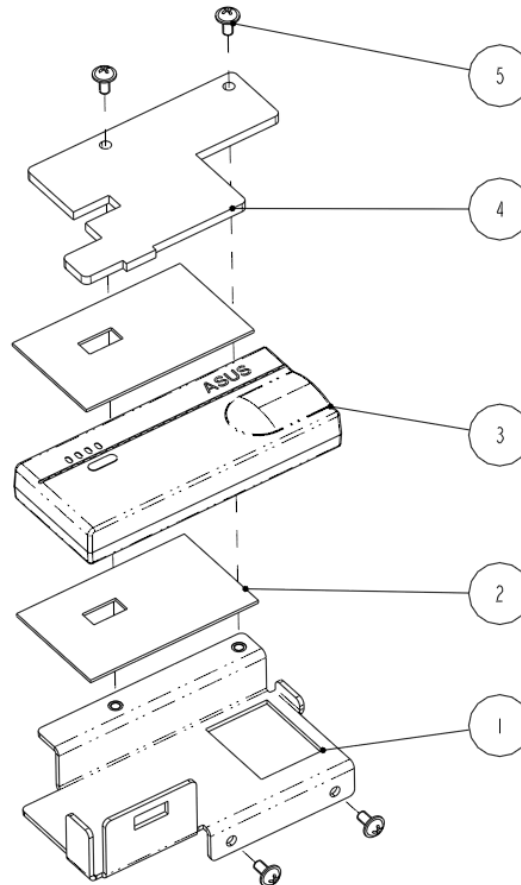


Parts List

SN	P/N	Description	Qty
1	047-000731-00	Cover overlay	1
2	6802-20-66725 or 043-004098-00	Card cover	1
3	6802-20-66726	CF card board fixture	1
4	M6T-030005---	Leaf, 178S30(1EA=406mm)	2
5	M04-002505---	Crosshead screw M3×6	2

6.4.10 6802 Internal Wireless AP Assembly

Exploded View



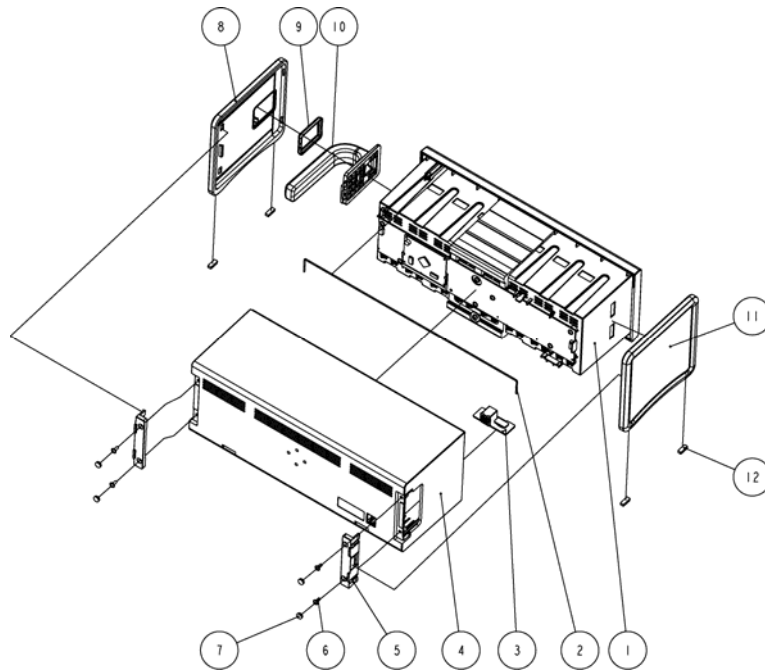
Parts List

SN	P/N	Description	Qty
1	042-000600-01	Wireless AP holder for T5	1
2	047-008023-00	Thermal pad for wireless card	2
3	023-000505-00	Wireless router 150Mbps Wi-Fi	1
4	042-007522-00	Cover for wireless LAN (T5)	1
5	M04-004012---	Cross pan head screw with washer M3X6	4
6	009-002895-00	Cable for AP wireless (6100)	1

6.5 SMR

6.5.1 SMR Assembly

Exploded View



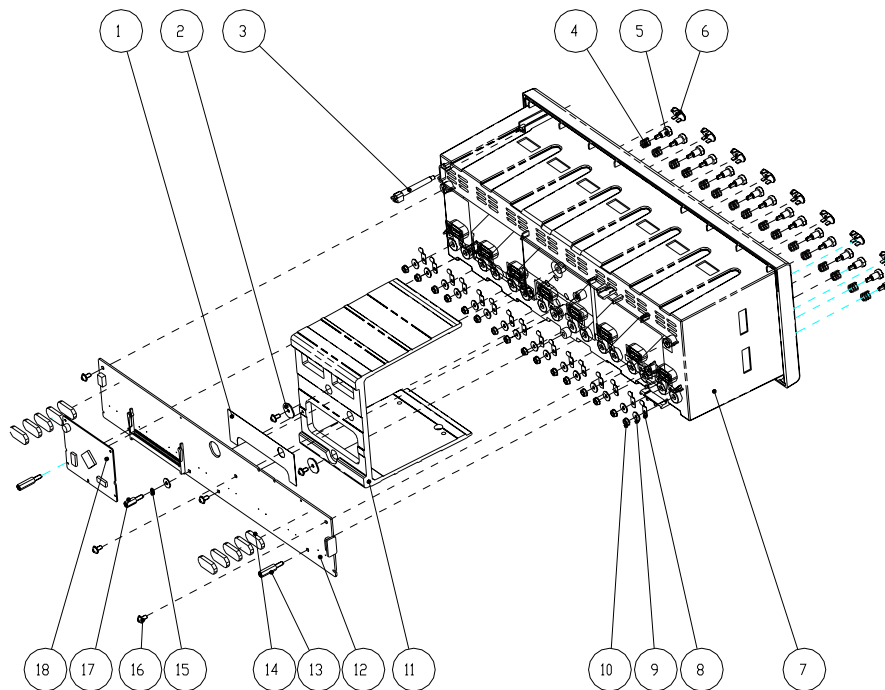
Parts List

SN	P/N	Description	Qty
1	6800-20-50249-51 or 6800-20-50249-52 or 043-004118-00 or 043-004119-00	Inside assembly	1
2	M6G-020015---	Hose	1
3	6800-30-51154	Interface board	1
4	6800-20-50250 or 043-004120-00	Rear housing	1
5	6800-20-50254 or 043-004054-00	Side plate fixture	2
6	M04-004015---	Crosshead screw M3×8	4
7	6800-20-50308 or 049-000650-00	Screw cap 2	4
8	6800-20-50253 or	Side plate, right	1

	043-004053-00		
9	6800-20-50262 or 049-000651-00	Rubber washer	1
10	6800-20-50251 or 043-004051-00	Handle	1
11	6800-20-50252 or 043-004052-00	Side plate, left	1
12	6800-20-50263	Cushion	4

6.5.2 SMR Inside Assembly

Exploded View



Parts List

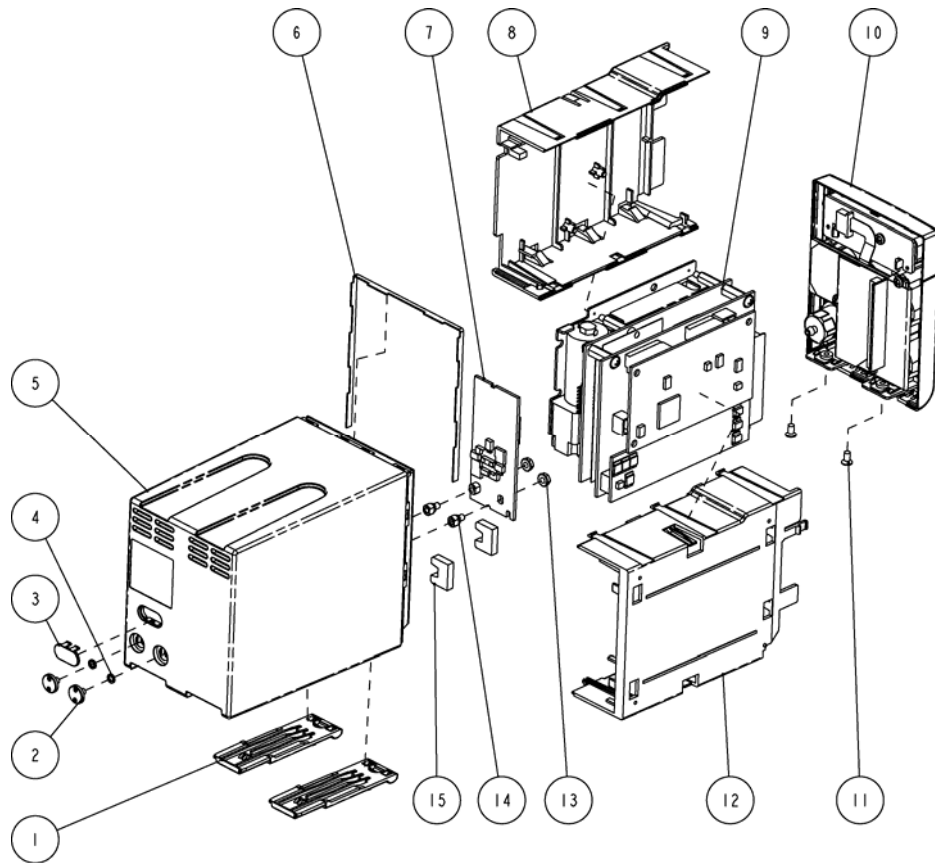
SN	P/N	Description	Qty
1	6800-20-50519	Insulating plate	1
2	9201-20-36010	Torsion spring washer	2
3	6800-20-50256	Light conducting pole	1
4	6800-20-50261	Contact spring	16
5	6800-21-51100	Contact screw	16

SN	P/N	Description	Qty
6	6800-20-50279	Infrared light filter	8
7	6800-20-50249 or 043-004118-00	SMR inside assembly	1
8	6800-20-50388	Leaf	16
9	M04-021024---	Washer	16
10	M04-011002---	Hex nut assy	16
11	6800-20-50257-51	Reinforced section steel	1
12	6800-30-51153	8-slot module rack communication board	1
13	M04-004306---	stub screw M3X16+8-8	2
14	6800-20-50233	Rubber feet	10
15	M04-000104---	spring washer GB93 3	1
16	M04-004012---	Crosshead screw M3×6	6
17	M04-000306---	stud screw M3X10+8-8, coated with antirust nickel	1
18	6800-30-50075	Nios II module	1

6.6 Parameter Modules

6.6.1 MPM Module

Exploded View



Parts List

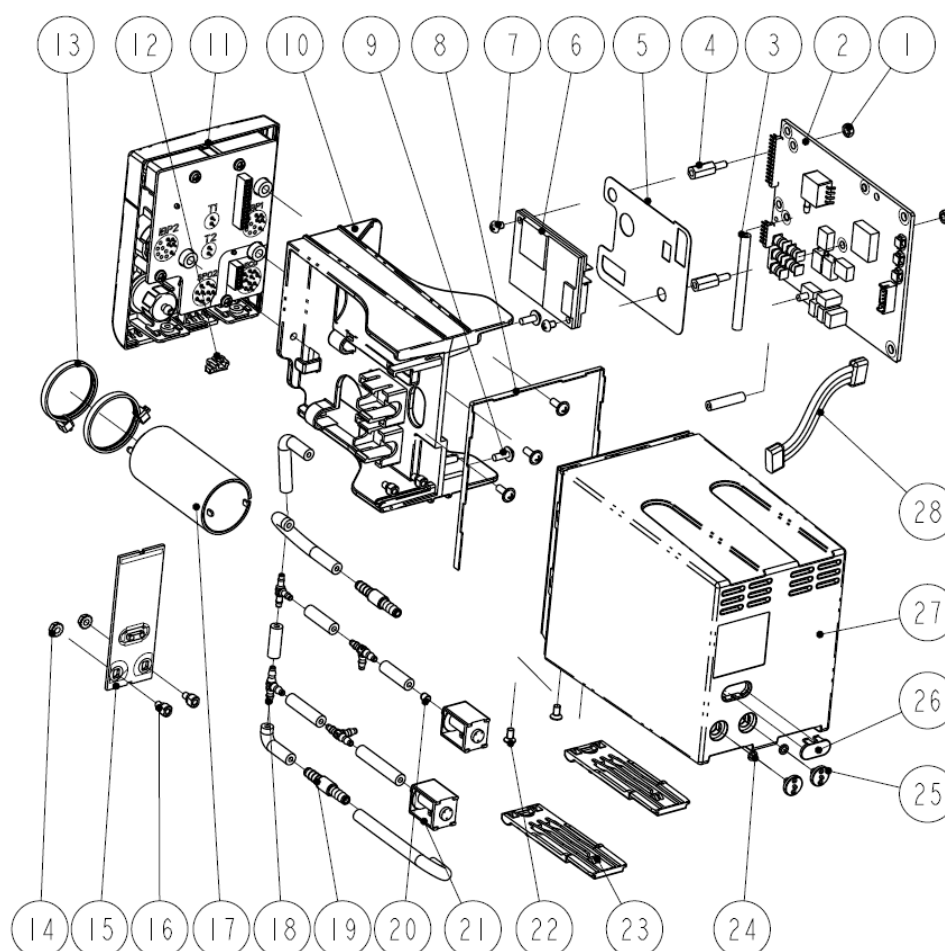
SN	P/N	Description	Qty
1	6800-20-50278 or 043-004037-00	Snap plate	2
2	6800-20-50293	Screw	2
3	6800-20-50279	Infrared light filter	1
4	M04-000104---	Spring washer	2
5	6800-20-50273 or 043-004031-00	2-slot module rear housing	1
6	6800-20-50339	Waterproof seal 02	1
7	6800-30-50684	MPM infrared communication backboard	1
8	6800-20-50275 or 043-004033-00	2-slot module inner housing, right side	1

9	M51A-30-80924	NIBP&MPM analog assembly	1
10	M51A-30-80870	MPM front housing assembly	1
11	M04-005005---	Crosshead screw M3×6	2
12	6800-20-50274 or 043-004032-00	2-slot module inner housing, left side	1
13	M04-011002---	M3 nut with spring washer	2
14	6800-20-50292	Fixer	2
15	6800-20-50870	Cushion	2

6.6.2 New MPM Module

New MPM module applies the integrative parameter board (ECG ASIC).

Exploded View



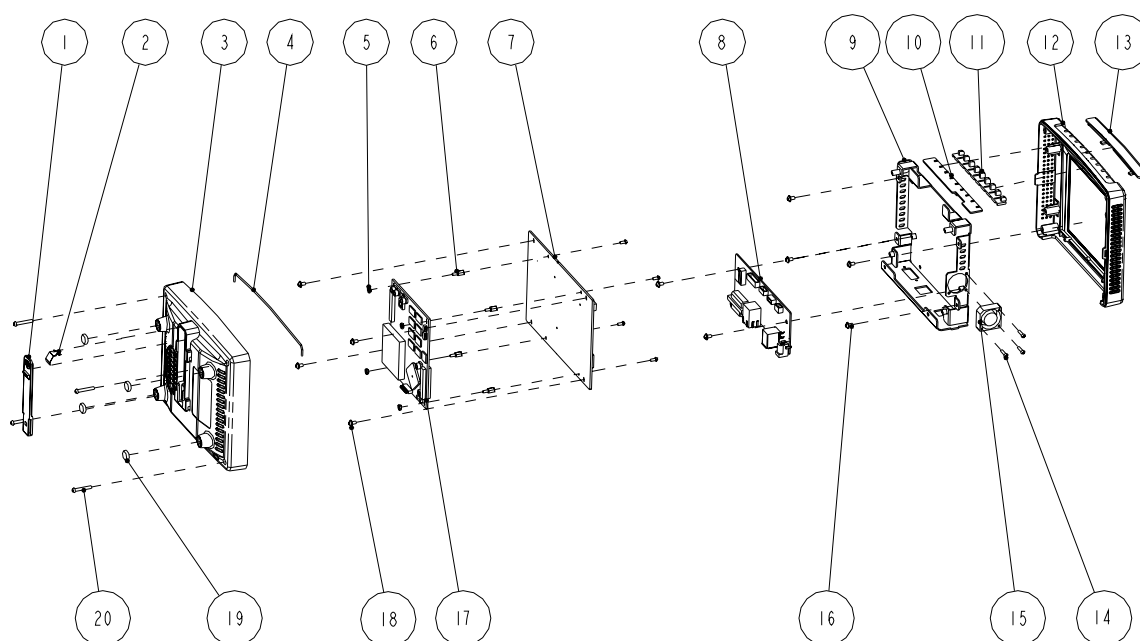
Parts List

SN	P/N	Description	Qty
1	M90-000156---	Plastic hex nut M3	2
2	051-000976-00	M51A Multi-parameter module, 5-lead, standard	1 As configured
	051-000977-00	M51A Multi-parameter module, 5-lead, full	
	051-000978-00	M51A Multi-parameter module, 12-lead, full	
	051-001037-00	M51A Multi-parameter module, 5-lead, full, IBP, Masimo	
3	A21-000007---	Silicone tube	1.65 inch
4	M90-000155---	Plastic hex stud screw, M3×12	2
5	047-006240-00	Insulation sheet for SpO ₂ board	1
6	051-000943-00	9008 V2.0 SpO ₂ board	1 As configured
	0671-00-0102-01	Nellcor SpO ₂ board	
	040-001149-00	Masimo, MS-2013, SpO ₂ board	
7	M04-051001-01	Crosshead screw M3×4	2
8	6800-20-50339	Waterproof seal 02	1
9	M04-004015---	Cross pan head screw with washer M3×8	5
10	043-001964-00 or 043-001964-01	Holder	1
11	115-011209-00	New MPM front panel assembly, Mindray SpO ₂	1 As configured
	115-011210-00	New MPM front panel assembly, Nellcor SpO ₂	
	15-011211-00	New MPM front panel assembly, Masimo2013 SpO ₂	
	115-011212-00	New MPM front panel assembly, Mindray SpO ₂ , no IBP	
	115-011213-00	New MPM front panel assembly, Nellcor SpO ₂ , no IBP	
	115-011214-00	New MPM front panel assembly, Masimo SpO ₂ , no IBP	
12	6800-20-50394 or 043-004039-00	Snap close	1
13	A90-000031---	Cable tie, CHS-4×150mm	2
14	M04-011002---	M3 nut with washer	2
15	115-011562-00	New M51A Infrared communication board	1

SN	P/N	Description	Qty
	115-011563-00	New M51A Infrared communication board, no IBP	As configured
16	6800-20-50292	Fixer	2
17	801-6800-00211-00	NIBP pump for M51A and iPM	1
18	S1-0103-00-0511	Tee connector, White, Nylon	4
19	0010-10-12408	Inline Filter	2
20	082-000098-00	630F Reducer	1
21	082-000864-00	Valve	1
22	M04-005005---	Flat head screw, Phillips M3×6	2
23	6800-20-50278 or 043-004037-00	Spanner	2
24	M04-000104---	Spring Washer	2
25	6800-20-50293	Screw	2
26	6800-20-50279	Infrared light filter	1
27	6800-20-50273 or 043-004031-00	Rear cover	1
28	009-002220-00	Cable for infrared Communication board	1

6.7 Remote Display Box

Exploded View

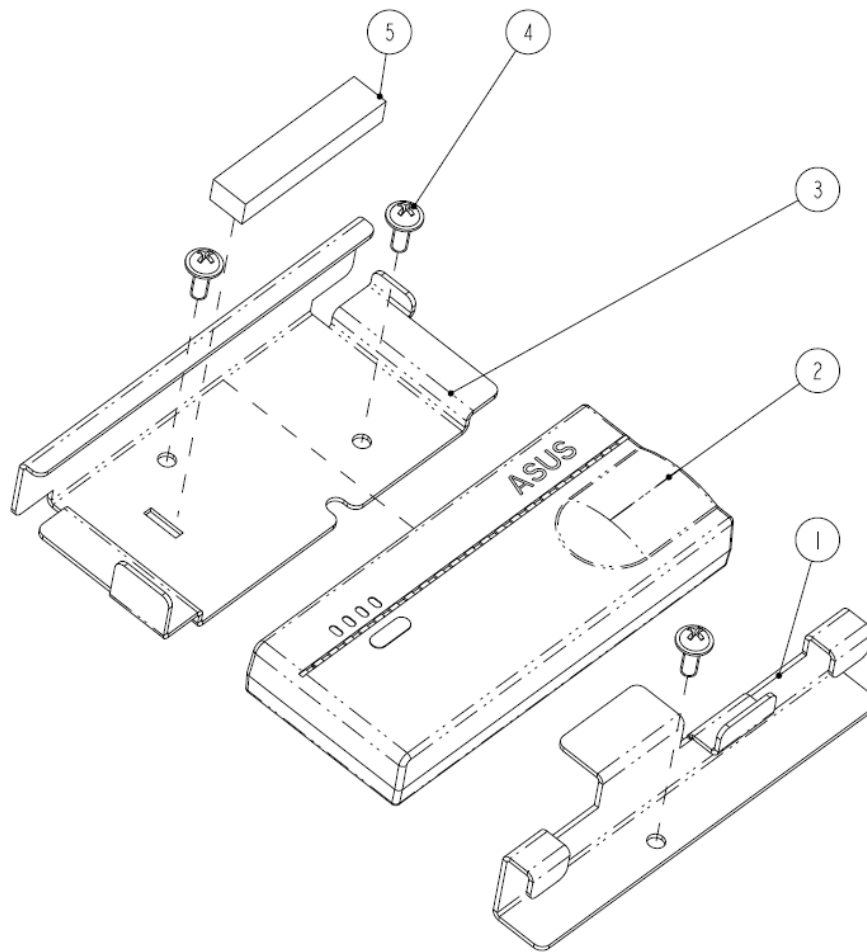


Parts List

SN	P/N	Description	Qty
1	M12A-20-75103	CIS lock catch	1
2	M12A-20-75107	Lock catch spring	1
3	M12A-21-75110	CIS bottom housing	1
4	M6G-020015---	Silicone tube	1
5	M04-011001---	Hex nut GB/T6170-2000M2.5	4
6	M04-060021---	Stud screw M2.5×7+6-6	4
7	M11A-30-75026	Remote display mother board	1
8	M11A-30-75000	Equipment interface board	1
9	M11A-20-75004-51	Remote display box rack	1
10	M12A-30-75002	Indicator board	1
11	M12A-20-75106	Light block	1
12	M12A-20-75101-51	CIS top housing	1
13	M12A-20-75102-52	Remote display box light cover	1
14	M04-004705---	Cross pan head screw M2.5×12	3
15	6800-20-50164	Module fan and cable	1
16	M04-004012---	Cross pan head screw with washer M3×6	10
17	6800-30-50058	MCF547x CPU module (basic configuration/lead-free process)	1
18	M04-051045---	Cross pan head screw M2.5×6	4
19	M12A-20-75105	CIS box cushion	4
20	M04-000505---	Cross pan head screw M3×20	4

6.8 New Wireless AP Package (ASUS) for 6800/6802

Exploded View



Part List

SN	P/N	Description	Qty
1	042-007507-00	Wireless AP/CF card mount	1
2	023-000505-00	Wireless router 150Mbps Wi-Fi	1
3	042-007506-00	Mounting plate for wireless LAN (ASUS)	1
4	M04-004015---	Cross recessed small pan head screw with washer M3X8	3
5	9201-20-36109	Wireless network card spacer	1
6	9201-20-53698	Network cable	1

6.9 Replaceable Parts

To replace the parts, please refer to *5 Repair and Disassembly* and the exploded views above.

NOTE

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- Here we list most replaceable parts. If you need more parts, please contact our Customer Service Department.
-

6.9.1 Main Unit

P/N	Description	Qty
Front housing assembly		
0000-10-11020	Backlight board	1
022-000014-00 or 022-000001-00	TPI inverter	1
0000-10-10752	LCD screen (SHARP)	1
0000-10-42633 (optional)	LCD screen (AU)	1
0000-10-10799	Touchscreen 12.1"	1
0000-10-10749 (optional)	Touchscreen 12.1"	1
6800-30-50082	Touchscreen control board	1
6800-30-51095 (optional)	Touchscreen control board (should be used in connection with the optional touchscreen 0000-10-10749)	1
6802-30-66812	Button (AU 12.1" screen G121SN01+anti-glare screen)	
6802-30-66814	Button (AU 12.1" screen G121SN01+touchscreen)	
6802-30-66682	Alarm LED board	1
6800-30-51132	Encoder assembly	
6800-30-50520	Encoder assembly	1
6802-30-66680	Power switch & LED board	1
6802-20-66732	Connecting cable between converter and keypad (Sharp)	1
6802-20-66731	12" SHARP screen connecting cable	1
6802-20-66673	Cable connecting the inverter and the button board	1
6802-20-66650	Flat cable connecting the LCD and the mother board	1
6802-20-66670	Cable connecting the alarm LED board and the button board	1

6802-20-66671	Cable connecting the power switch & LED board and button board	1
6802-20-66672	Cable between the touchscreen control board and the button board	1
6802-30-66890	DPM6 service kit of anti-glare screen front housing	1
6802-30-66891	DPM6 service kit of touchscreen front housing	1
6812-30-66986	DPM6(6812) service kit of anti-glare screen front housing	1
6812-30-66987	DPM6(6812) service kit of touchscreen front housing	1
Main unit		
6006-20-39435	Recorder cover	1
6802-20-66665	Cable connecting the recorder and the mother board	1
6802-20-66666	Cable connecting the CF card and the mother board	1
6802-20-66684	Cable connecting the button board and the mother board	1
043-000059-00 or 043-000059-01 or *6812-20-67000-51 or 043-004115-00 or 043-000472-00	Interface board cover	1
6802-20-66717	Main support	1
6802-20-66718	Recorder fixing plate	1
6802-30-66763	Battery compartment assembly	1
6802-30-66653	Battery interface board	1
6802-20-66799	Cable connecting the Integral module rack and the mother board	1
6802-30-66767 or *6812-30-66993 or 115-003231-00	Master control board assembly	1
6802-30-66769 or *6812-30-66994 or 115-003230-00	Interface assembly	1
115-001906-00 or *6812-30-66995	6802 CF card assembly (211 driving board) CF card slot cover kit	1
6802-30-66768	Fan assembly	1
6802-20-66668	Speaker and cables	1
TR6F-30-67306 or 115-001290-00	TR6F recorder	1
6802-30-66770	Power module assembly	1

6802-30-66651	Power board	1
6800-30-50073	MPM module rack communication board	1
6800-30-50075	NiosII module	1
6802-20-66669	Cable connecting the main unit and the infrared communication board	1
051-000244-00	6802 3-slot module rack communication board	1
6800-20-50198-51	Key pad	1
6802-30-66766	Rear housing assembly	1
6800-20-50672	Cable connecting the main board and the DIV interface board	1
6800-20-50673	Cable connecting the main board and the USB_Hub board	1
6802-20-66664	Cable connecting the DVI interface board and the mother board	1
6802-20-66675	Cable connecting USB_Hub board and the mother board	1
6802-30-66659 or *6812-30-66999 or 051-000020-00	USB Hub board 6812 network interface board USB interface board (no analog output)	1
6802-30-66657 or 051-000019-00 or 051-000019-01	DVI interface board or DVI interface board (can support IABP) or DVI interface board (standard configuration)	1

6.9.2 SMR

P/N	Description	Qty
6800-30-50075	NiosII module	1
6800-30-50078	SMR communication board	1
6800-30-50080	SMR interface board	1
6800-30-50667	SMR indicating lamp assembly	1
6800-20-50249-51	SMR inside assembly	1
6800-20-50250-51	SMR rear housing	1
6800-20-50251	SMR handle	1
6800-20-50319	Cable connecting the SMR and the main unit	
6800-21-50167	SMR indicating lamp cable	

6.9.3 Parameter Modules

P/N	description	Qty
6800-30-50486	BIS Module	1
6800-30-50488	RM module	1
6800-30-50487	CAPNOSTAT CO ₂ module	1
6800-30-50559	ORIDION CO ₂ module	1
6800-30-50500	Mindray sidestream CO ₂ module (M02B)	1
115-020189-00	Mindray sidestream CO ₂ Module (M02C)	1
6800-30-50491	ICG module	1
6800-30-50501	AG module (with O ₂ /BIS)	1
6800-30-50502	AG module (with O ₂)	1
6800-30-50503	AG module (without O ₂ /BIS)	1
6800-30-50504	AG module (without O ₂)	1
6800-30-50505	Manual AG module (with O ₂ /BIS)	1
6800-30-50506	Manual AG module (with O ₂)	1
6800-30-50507	Manual AG module (without O ₂ /BIS)	1
6800-30-50508	Manual AG module (without O ₂)	1
6800-30-50484	C.O. module	1
6800-30-50485	IBP module	1
115-007270-00	PiCCO module	1
115-007273-00	ScvO ₂ module	1
115-007276-00	BeneLink module	1
115-018152-00	EEG module	1
115-018518-00	NMT module	1
115-007277-00	BeneLink ID adapter	1
M51A-30-80873	MPM-1 module(Mindray SpO ₂ /3/5 lead)	1
115-004900-00	MPM-2 module(MasimoSpO ₂ /3/5 lead)	1
M51A-30-80879	MPM-3 module(NellcorSpO ₂ /3/5 lead)	1
M51A-30-80880	MPM-4 module (MindraySpO ₂ /12 lead)	1
115-004904-00	MPM-5 module(MasimoSpO ₂ /12 lead)	1
M51A-30-80886	MPM-6 module(NellcorSpO ₂ /12 lead)	1
M51A-30-80900	MPM-7 module main unit(MindraySpO ₂ /3/5 lead/without IBP)	1
115-004909-00	MPM-8 module(Masimo SpO ₂ /3/5 lead/without IBP)	1
M51A-30-80906	MPM-9 module(Nellcor SpO ₂ /3/5 lead/without IBP)	1

P/N	description	Qty
M51A-30-80907	MPM-10 module main unit (MindraySpO ₂ /12 lead/without IBP)	1
115-004912-00	MPM-11 module (Masi SpO ₂ /12 lead /without IBP)	1
M51A-30-80913	MPM-12 module (Nell SpO ₂ /12 lead/without IBP)	1
115-010731-00	New MPM2 module(Masimo SpO ₂ /3/5lead)	1
115-010734-00	New MPM5 module(Masimo SpO ₂ /12lead)	1
115-010738-00	New MPM8 module(Masimo SpO ₂ /3/5lead/no IBP)	1
115-010740-00	New MPM11 module(Masimo SpO ₂ /12lead/no IBP)	1
115-010744-00	New MPM3 module(Nellcor SpO ₂ /3/5lead)	1
115-010747-00	New MPM6 module(Nellcor SpO ₂ /12lead)	1
115-010750-00	New MPM9 module(Nellcor SpO ₂ /3/5lead/no IBP)	1
115-010753-00	New MPM12 module(Nellcor SpO ₂ /12 lead)	1
115-010755-00	New MPM1 module(Mindray SpO ₂ /3/5 lead)	1
115-010758-00	New MPM4 module(Mindray SpO ₂ /12 lead)	1
115-010761-00	New MPM7 module(Mindray SpO ₂ /3/5lead/no IBP)	1
115-010764-00	New MPM10 module(Mindray SpO ₂ /12lead/no IBP)	1
6800-30-50128	Mindray SpO ₂ flexible cable kit	
6800-30-50132	Nellcor SpO ₂ flexible cable kit	
6800-30-50130	Masimo SpO ₂ flexible cable kit	
6800-20-50662	Air pump cable	
6800-20-50663	Fast-release valve cable	
6800-20-50664	Slow-release valve cable	
6800-20-50674	Cable from ICG module to infrared communication board	
6800-20-50683	Mindray CO ₂ infrared communication cable	
6800-20-50306	AG module cable	
6800-20-50309	ICG module communication cable	
6800-21-50310	BIS interface cable	
6800-21-50311	CO interface cable	
6800-21-50312	IBP interface cable	
6800-21-50313	Mainstream CO ₂ interface cable	
6800-20-50315	ICG power cord	
6800-20-50316	RM infrared detection board cable	
6800-20-50160	Module button board cable	
040-000125-00	Patient Interface Cable (BIS module service part)	
040-000674-00	For service only, BISx Kit (186-0199-MR)	
040-000675-00	For service only, BISx4 Kit (186-1030-MR)	

P/N	description	Qty
040-000676-00	For service only, BISx Host Cable (186-0201-MR)	
009-001770-00	RJ45 connecting cable	
009-001767-00	Serial port adapting cable, type A	
009-001768-00	Serial port adapting cable, type B	
009-001769-00	Serial port adapting cable, type C	
009-001765-00	Cable, Infrared Board to Interface Board	
009-001254-00	AP&CVP socket with signal cable	
009-001253-00	PiCCO socket with signal cable	
009-001255-00	ScvO ₂ socket with signal cable	
009-002220-00	Cable for infrared Communication board	
009-003211-00	NMT signal cable	
100-000070-00	EEG signal cable	

7 Upgrade

7.1 Introduction

You can upgrade parameter modules, functional assemblies and software by connecting the patient monitor to a PC running the System Update Tool.

NOTE

- **If you have to disassemble the patient monitor for software upgrade, be sure to eliminate the static charges before disassembling the equipment. When disassembling the part labeled with a ESD warning symbol, make sure you are wearing electrostatic discharge protection such as antistatic wristband or gloves to avoid damaging the equipment.**
 - **Properly connect and route the cables and wires when reassembling the equipment to avoid short circuit.**
 - **Select appropriate screws to assemble the equipment. If unfit screws are tightened by force, the equipment may be damaged and the screws or part may fall off during use, causing unpredictable equipment damage or human injury.**
 - **Follow correct sequence to disassembly the equipment. Otherwise, the equipment may be damaged permanently.**
 - **Disconnect all the cables before disassembling any parts. Be careful not to damage any cables or connectors.**
 - **Properly place removed screws and disassembled parts, preventing them from being lost or contaminated.**
-

7.2 Upgrading Parameter Modules

You can upgrade the following parameter modules:

Parameter module	PN	Description	Remark
MPM module	M51A-30-80873	MPM-1Module(MINDRAYSpO ₂ , 3/5lead)	/
	115-004900-00	MPM-2 Module (Masimo MS-2011, 3/5lead)	/
	M51A-30-80879	MPM-3 Module (NellcorSpO ₂ , 3/5 lead)	/
	M51A-30-80880	MPM-4 Module (MINDRAYSpO ₂ , 12 lead)	/
	115-004904-00	MPM-5 Module (Masimo MS-2011, 12 lead)	/
	M51A-30-80886	MPM-6 Module (NellcorSpO ₂ , 12 lead)	/
	M51A-30-80900	MPM-7Module(MindraySpO ₂ , 3/5 lead, no IBP)	/
	115-004909-00	MPM-8Module(Masimo MS-2011, 3/5 lead, no IBP)	/
	M51A-30-80906	MPM-9Module(NellrSpO ₂ , 3/5 lead, no IBP)	/
	M51A-30-80907	MPM-10Module(MindraySpO ₂ , 12 lead, no IBP)	/
	115-004912-00	MPM-11Module(Masimo MS-2011, 12 lead , no IBP)	/
	M51A-30-80913	MPM-12Module(NellSpO ₂ , 12 lead, no IBP)	/
New MPM module	115-010731-00	MPM2 module(Masimo SpO ₂ , 3/5lead)	
	115-010734-00	MPM5 module(Masimo SpO ₂ , 12lead)	
	115-010738-00	MPM8 module(Masimo SpO ₂ , 3/5lead, no IBP)	
	115-010740-00	MPM11 module(Masimo SpO ₂ , 12lead, no IBP)	
	115-010744-00	MPM3 module(Nellcor SpO ₂ , 3/5lead)	
	115-010747-00	MPM6 module(Nellcor SpO ₂ , 12lead)	
	115-010750-00	MPM9 module(Nellcor SpO ₂ , 3/5lead, no IBP)	
	115-010753-00	MPM12 module(Nellcor SpO ₂ , 12 lead)	
	115-010755-00	MPM1 module(Mindray SpO ₂ , 3/5lead)	
	115-010758-00	MPM4 module(Mindray SpO ₂ , 12lead)	
	115-010761-00	MPM7 module(Mindray SpO ₂ , 3/5lead, no IBP)	
	115-010764-00	MPM10 module(Mindray SpO ₂ , 12lead, no ibp)	

IBP module	6800-30-50850	IBP module upgrade package (without accessories)	/
C.O. module	6800-30-50849	C.O. module upgrade package (without accessories)	/
CO ₂ module	6800-30-50139	M02B CO ₂ module upgrade package (for adult and pediatric patients, with accessories)	Sidestream (2 slots)
	6800-30-50141	M02B CO ₂ module upgrade package (for neonatal patient, with accessories)	Sidestream (2 slots)
	115-020189-00	CO ₂ module	Sidestream (1 slot)
	6800-30-50820	Oridion CO ₂ module upgrade package (with accessories)	Microstream
	6800-30-50852	Capnostat CO ₂ module upgrade package (with accessories)	Mainstream
AG module	6800-30-50841	AG module upgrade package (with O ₂ , BIS, and accessories)	/
	6800-30-50842	AG module upgrade package (with O ₂ and accessories)	/
	6800-30-50843	AG module upgrade package (without O ₂ , with BIS and accessories)	/
	6800-30-50844	AG module upgrade package (without O ₂ , with accessories)	/
ICG module	6800-30-50854	ICG module upgrade package (with accessories)	/
BIS module	6800-30-50427	BIS module upgrade package (for pediatric patients, with accessories)	/
	6800-30-50880	BIS module upgrade package (with accessories)	/
RM module	6800-30-50853	RM module upgrade package (with accessories)	/
CCO/SvO ₂ module	801-6800-00104-00	CCO/SvO ₂ module upgrade package	/
PiCCO Module	115-007589-00	PiCCO function upgrade kit	/
ScvO ₂ Module	115-007590-00	ScvO ₂ function upgrade kit	/
EEG module	115-018152-00	EEG module	/
NMT module	115-018518-00	NMT module	/

You can plug and unplug parameter modules during patient monitoring.

Refer to the Operator's Manual for the use of parameter modules.

7.3 Upgrading Functional Assemblies

You can upgrade the following functional assemblies:

Functional assembly	PN	Description	Remark
SMR	6800-30-50641	SMR kit	/
Wireless network	801-6802-00030-00	6802 wireless network upgrade kit	Internal AP, for standard- configured patient monitor
	801-6802-00031-00	6802 wireless network upgrade kit	Internal AP, for fully configured patient monitor
	801-6800-00002-00	Wireless network adaptor kit	External AP
Recorder	TR6F-30-67318	Recorder upgrade kit	/
Analog output	801-6802-00006-00	DVI interface board (FRU)	/

The patient monitor can be connected to network through wireless AP. Authorized personnel are required to connect and set up the wireless network, and then carry out the performance test.

7.3.1 Upgrading SMR

The SMR can be connected to the patient monitor through SMR connector via a SMR cable.

Refer to the Operator's Manual for detail.

7.3.2 Upgrading Wireless Network Function

Refer to corresponding section to install the wireless network assembly in your patient monitor.

Refer to the Operator's Manual for the use of wireless network.

7.3.3 Upgrading Recorder

Refer to corresponding section to install the recorder in your patient monitor.

Refer to the Operator's Manual for the use of recorder.

7.3.4 Upgrading Analog Output

Refer to corresponding section to replace the interface board assembly and DVI interface board.

Refer to the Operator's Manual for use of the analog output.

7.3.5 Upgrading CF storage card function

Refer to *Removing the CF Card Assembly* section to install CF card kit. Then insert the CF card into the slot.

Refer to the Operator's Manual for the use of CF card.

NOTE

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- **When upgrading wireless network, analog output and CIS function for a patient monitor with standard configuration, you have to replace old PCBAs in the patient monitor with corresponding PCBAs included in the upgrade kit and remove the covers of related connectors.**
-

7.4 Upgrading Software

You can upgrade the software with the System Update Tool (PN: 110-000493-00) through network.. This tool can directly run on a PC. Through network or by connecting the patient monitor to a PC via a crossover network cable, you can upgrade the following programs:

Software	PN	Description	Remark
Boot program	6800-30-50770	BIOS program	Bootstrap program for monitor main control unit, implementing hardware drive and initialization For CE
	110-001594-00	BIOS program	For FDA
System software	6800-30-50701	System software	FDA
	110-000414-00	System software	CE
Multilingual library	6800-30-50702	Language library	FDA
	110-000421-00	Language library	CE
BMP resource file	6800-30-50767	Icon library	FDA
	110-000422-00	Icon library	CE
	110-000807-00	Startup Logo	CE
	6802-30-66876	Startup Logo	FDA
FPGA program	110-000815-00	FPGA configuration chip software for mother board	CE
	6802-30-66895	FPGA configuration chip software for mother board (MDDR)	FDA
Audio data file	6800-30-50872	Audio data software for mother board	FDA
	110-001423-00	Audio data software for mother board	CE
Module rack program	6800-30-50863	FPGA configuration chip software for Module rack NIOS board	/
MPM module	0010-30-43083	BIOS program for parameter module(BF531)	The System Update Tool for MPM

program	M51A-30-80950	DSP board M25P16 software for M51A module	
	M51A-30-80951	DSP board LPC2131 software for M51A module	
	M51A-30-80952	7024 software for M51A module ECG board	
	110-000539-00	SpO ₂ module software for 9008	
	M51A-30-80953	7024 software for M51A module SpO ₂ board	
	M51A-30-80956	M25P16 software for M51A module DSP board, Mortara algorithm	
New MPM module program	110-001979-00	BIOS program for parameter module(BF512)	The System Update Tool for M51A_2
	110-001839-00	M51A V2.0 DSP(BF512) software	
	110-001978-00	M51A V2.0 DSP(Mortara) software	
	110-001842-00	9008 V2.0 SpO ₂ software	

7.4.1 How to Upgrade Software

Before software upgrade, check software version by selecting [**Main Menu**] → [**Maintenance>>**] → [**Factory Maintenance>>**] → enter the required password → [**Software Version>>**]. If your BIOS program has a version prior to 2.0, upgrade it to version 2.0 or greater before upgrading other programs. If you use System Update Tool (PN: 110-000493-00) version 2.0 or greater, make sure that BIOS program version is 2.0 or greater before upgrade. Please refer to the following procedure to upgrade the software:

1. Connect the patient monitor to a PC through a crossover network cable. Set the IP to 77.77.1.XX, the subnet mask to 255.255.255.0.
2. Run the System Update Tool in the PC. Select [**BeneView**] in the [**Machine Type Selection**] window and confirm.
3. Select [**Select Package**] in the main screen of the Mindray Patient Monitor Software Upgrade Tool. Choose the package you want to upgrade and click [**Start**].
4. Start up the patient monitor and press quickly the [**Silence**]+[**Main Menu**] hardkeys to enter upgrade mode. Determine if the software is upgraded successfully according to the prompt message of the patient monitor when finishing upgrading the software.

If your BIOS program has a version prior to 5.0(CE) or 4.2.0(FDA),, you should press [**Silence**] hardkey before you start the patient monitor, and do not release it until you enter the upgrade screen.

Software upgrade is taken in the sequence of BIOS program → system software package (including system software, language library, and BMP resource files) → FPGA package (including mother board FPGA software and audio data files) → module rack FPGA package → MPM module software.

7.4.1.1 Upgrading BIOS program

If your BIOS program is version 1.2 or prior, the patient monitor has a blank screen during software upgrade. For BIOS program version greater than 1.2, the patient monitor enters the upgrade screen and presents related prompts during software upgrade.

For example, to upgrade BIOS program from version 2.0 to 3.0 or greater,

1. Run the System Update Tool, version 3.0.
2. Download the BIOS upgrade program (110-000296-00). Create a system package using the BIOS upgrade program and download it to the patient monitor. Turn off the patient monitor after you finish downloading the package.
3. Select BIOS package version 3.0 or greater, and then select the “Start” button on the System Update Tool screen. Turn on the patient monitor by pressing the power switch (you are not allowed to press other buttons at same time). Thus you can upgrade the BIOS program to version 3.0.

It should be noted that the patient monitor shows a blank screen for about 20 seconds when you turn on the patient monitor for the first time after successful software upgrade. This is because the new BIOS program is restoring the master boot from backup BIOS program.

7.4.1.2 Upgrading System Software

To upgrade system software version 02.02.00 or prior, BIOS program version 2.0 or greater and System Update Tool version 2.0 or greater are required.

The language library and icon library shall be upgraded at same time when system software is upgraded.

7.4.1.3 Upgrading FPGA Program

If your FPGA program is version 1.0, first you have to upgrade it to version 2.0 using the System Update Tool version 1.4, and then upgrade it to the latest version using the System Update Tool version 2.0 or greater.

Because two display memory models are available for the mother board, there are two mother board FGPA programs. The two FGPA programs can be created into one package to upgrade the FPGA programs for mother boards equipped with different display memories. The audio data file shall be upgraded at same time when FPGA program is upgraded.

For details, refer to the help and instructions included in the System Update Tool, or consult your service personnel.



CAUTION

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- **Disconnect the patient monitor from patient and make sure important data are saved before upgrade.**
 - **Do not shut down or power off the equipment when upgrading the BIOS program and FPGA program. Otherwise, the equipment may break down.**
 - **Program upgrade should be performed by qualified service personnel only.**
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NOTE

-
- **After upgrading the BIOS program, you have to upgrade system program and other programs to ensure compatibility.**
 - **Make sure the version of the upgrade package is you desired one. If you want to obtain the latest upgrade package, contact Mindray Customer Service Department.**
 - **To upgrade the MPM module from Mindray ECG algorithm to Mortara ECG algorithm, you have to send the MPM module to manufacture.**
-

A Electrical Safety Inspection

The following electrical safety tests are recommended as part of a comprehensive preventive maintenance program. They are a proven means of detecting abnormalities that, if undetected, could prove dangerous to either the patient or the operator. Additional tests may be required according to local regulations.

All tests can be performed using commercially available safety analyzer test equipment. These procedures assume the use of a 601PROXL International Safety Analyzer or equivalent safety analyzer. Other popular testers complying with IEC 60601-1 used in Europe, such as Fluke, Metron, or Gerb, may require modifications to the procedure. Please follow the instructions of the analyzer manufacturer.

The consistent use of a safety analyzer as a routine step in closing a repair or upgrade is emphasized as a mandatory step if an approved agency status is to be maintained. The safety analyzer also proves to be an excellent troubleshooting tool to detect abnormalities of line voltage and grounding, as well as total current loads.

A.1 Power Cord Plug

Test Item		Acceptance Criteria
The power plug	The power plug pins	No broken or bent pin. No discolored pins.
	The plug body	No physical damage to the plug body.
	The strain relief	No physical damage to the strain relief. No plug warmth for device in use.
	The power plug	No loose connections.
The power cord		No physical damage to the cord. No deterioration to the cord.
		For devices with detachable power cords, inspect the connection at the device.
		For devices with non-detachable power cords, inspect the strain relief at the device.

A.2 Device Enclosure and Accessories

A.2.1 Visual Inspection

Test Item	Acceptance Criteria
The enclosure and accessories	No physical damage to the enclosure and accessories.
	No physical damage to meters, switches, connectors, etc.
	No residue of fluid spillage (e.g., water, coffee, chemicals, etc.).
	No loose or missing parts (e.g., knobs, dials, terminals, etc.).

A.2.2 Contextual Inspection

Test Item	Acceptance Criteria
The enclosure and accessories	No unusual noises (e.g., a rattle inside the case).
	No unusual smells (e.g., burning or smoky smells, particularly from ventilation holes).
	No taped notes that may suggest device deficiencies or operator concerns.

A.3 Device Labeling

Check the labels provided by the manufacturer or the healthcare facility are present and legible.

- Main unit label
- Integrated warning labels

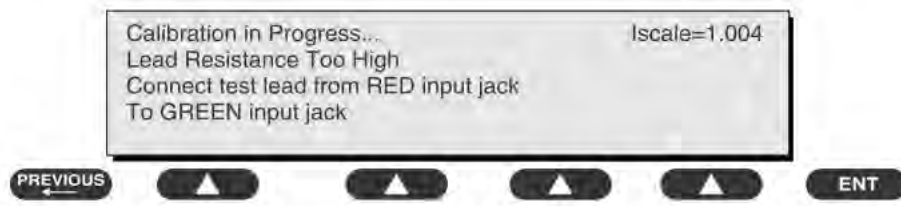
A.4 Protective Earth Resistance

Protective Earth Resistance is measured using the RED test lead attached to the DUT Protective Earth terminal or enclosure. Select the test current (25 amp) by pressing SOFT KEY 3. The front panel outlet power is turned off for this test.

The following conditions apply: L1 and L2 Open.

Preparation

1. First select the test current that will be used for performing the Protective Earth Resistance test by pressing AMPERES (SOFT KEY 3).
2. Connect the test lead(s) between the RED input jack and the GREEN input jack.
3. Press CAL LEADS. The 601PRO will measure the lead resistance, and if less than 0.150 Ohms, it will store the reading and subtract it from all earth resistance readings taken at the calibrated current.



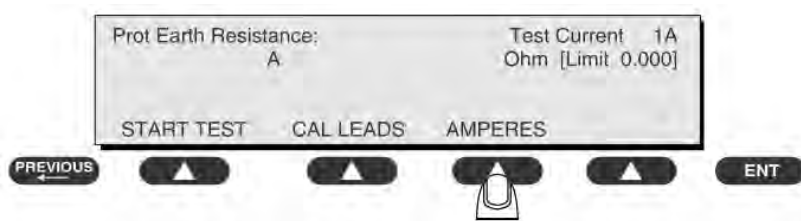
If the calibration fails, the previously stored readings will be used until a passing calibration has occurred.

WARNING

- **During Earth Resistance testing, the DUT must be plugged into the 601PRO front outlet. If the DUT fails Earth Resistance, discontinue tests and label the device defective.**
-

To Perform the Test

1. From the MAIN MENU, or with the outlet unpowered, plug the DUT into the 601PRO front panel outlet.
2. Attach the 601PRO RED input lead to the device's Protective Earth terminal or an exposed metal area.
3. Press shortcut key 3. The Protective Earth Resistance test is displayed.
4. Press SOFT KEY 3 to select a test current (25 amp). The selected test current is displayed in the upper right corner of the display.



5. Press START TEST to start the test. The test current is applied while resistance and current readings are taken. This takes approximately 5 seconds.
6. Press the print data key at any time to generate a printout of the latest measurement(s).

NOTE

- When "Over" is displayed for Ohms, this signifies that a valid measurement was not obtained because either an open connection was detected or that the measurement was not within range. Readings greater than 9.999 Ohms will be displayed as Over.
-

In Case of Failure

Once it reaches the limitation, stop using and inform the Customer Service Engineer for analysis and disposal.

LIMITS

ALL COUNTRIES R = 0.2 Ω Maximum

A.5 Earth Leakage Test

Run an Earth Leakage test on the device being tested before performing any other leakage tests.

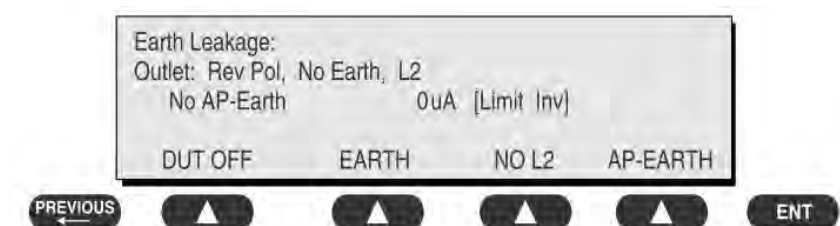
Leakage current is measured the following ways:

- Earth Leakage Current, leakage current measured through DUT outlet Earth
- Earth Leakage Current AP-EARTH (ALL Applied Parts connected to Earth), leakage current measured through DUT outlet Earth

There is no need to attach a test lead; the 601PRO automatically connects the measuring device internally.

To Perform the Test

1. From the MAIN MENU, or with the outlet unpowered, plug the DUT into the 601PRO front panel outlet, and turn on the device.
2. Attach the device's applied parts to the 601PRO applied part terminals if applicable.
3. Press shortcut key 4. The Earth Leakage test appears on the display, and the test begins immediately:



-
- SOFT KEY 1 toggles the DUT outlet Polarity from Normal to Off to Reverse.
 - SOFT KEY 2 toggles the DUT outlet from Earth to No Earth.
 - SOFT KEY 3 toggles the DUT outlet from L2 to No L2.
 - SOFT KEY 4 toggles the AP to Earth to No AP to Earth.
4. Press the print data key at any time to generate a printout of the latest measurement.

In Case of Failure

- Check any broken of the enclosure. Replace any defective part.
- Inspect wiring for bad crimps, poor connections, or damage.
- Test the wall outlet; verify it is grounded and is free of other wiring abnormalities. Notify the user or owner to correct any deviations. As a work around, check the other outlets to see if they could be used instead.
- Change another probe to confirm if the fail is caused by console.
- If the leakage current measurement tests fail on a new unit and if situation can not be corrected, submit a Safety Failure Report to document the system problem. Remove unit from operation.
- If all else fails, stop using and inform the Customer Service Engineer for analysis and disposal.

LIMITS

For UL60601-1,

- ◆ 300 μ A in Normal Condition
- ◆ 1000 μ A in Single Fault Condition

For IEC60601-1,

- ◆ 500 μ A in Normal Condition
- ◆ 1000 μ A in Single Fault Condition

A.6 Patient Leakage Current

Patient leakage currents are measured between a selected applied part and mains earth. All measurements have a true RMS only response.

Preparation

Perform a calibration from the Mains on Applied Part menu.

The following outlet conditions apply when performing this test:

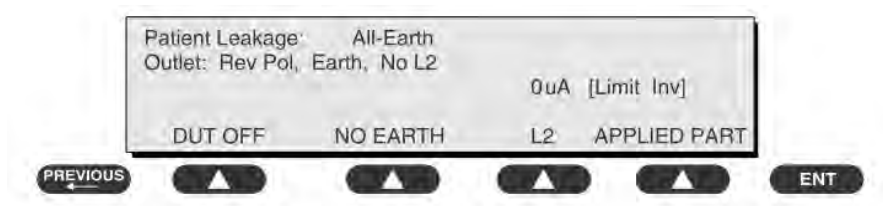
- Normal Polarity, Earth Open, Outlet ON Normal Polarity, Outlet ON
- Normal Polarity, L2 Open, Outlet ON Reversed Polarity, Outlet ON
- Reversed Polarity, Earth Open, Outlet ON Reversed Polarity, L2 Open, Outlet ON

WARNING

- **If all of the applied parts correspond to the instrument type, the applied parts will be tied together and one reading will be taken. If any of the applied parts differ from the instrument type, all applied parts will be tested individually, based on the type of applied part. This applies to Auto and Step modes only.**
-

To Perform the Test

1. From the MAIN MENU, or with the outlet unpowered, plug the DUT into the 601PRO front panel outlet, and turn on the device.
2. Attach the applied parts to the 601PRO's applied part terminals.
3. Press shortcut key 6. The Patient Leakage test is displayed, and the test begins immediately.



4. Press APPLIED PART (SOFT KEY 4) at any time to select the desired applied part leakage current.
5. Modify the configuration of the front panel outlet by pressing the appropriate SOFT KEY on the 601PRO.
6. Press the print data key at any time to generate a printout of the latest measurement.


In Case of Failure

- Check any broken of the enclosure. Replace any defective part.
- Inspect wiring for bad crimps, poor connections, or damage.
- Test the wall outlet; verify it is grounded and is free of other wiring abnormalities. Notify the user or owner to correct any deviations. As a work around, check the other outlets to see if they could be used instead.
- Change another probe to confirm if the fail is caused by console.
- If the leakage current measurement tests fail on a new unit and if situation can not be corrected, submit a Safety Failure Report to document the system problem. Remove unit from operation.
- If all else fails, stop using and inform the Customer Service Engineer for analysis and disposal.

LIMITS

For CF  applied parts

- ◆ 10 μ A in Normal Condition
- ◆ 50 μ A in Single Fault Condition

For BF  applied parts

- ◆ 100 μ A in Normal Condition
- ◆ 500 μ A in Single Fault Condition

A.7 Mains on Applied Part Leakage

The Mains on Applied Part test applies a test voltage, which is 110% of the mains voltage, through a limiting resistance, to selected applied part terminals. Current measurements are then taken between the selected applied part and earth. Measurements are taken with the test voltage (110% of mains) to applied parts in the normal and reverse polarity conditions as indicated on the display.

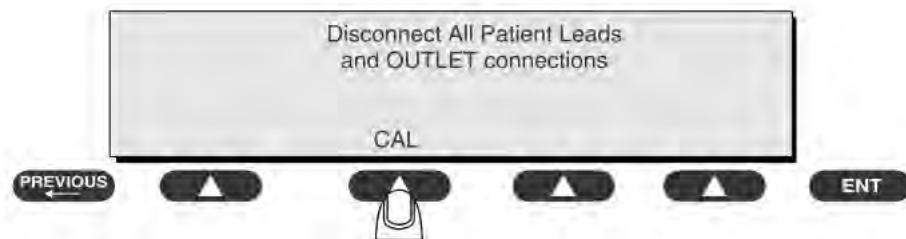
The following outlet conditions apply when performing the Mains on Applied Part test.

- Normal Polarity;
- Reversed Polarity

Preparation

To perform a calibration from the Mains on Applied Part test, press CAL (SOFT KEY 2).

1. Disconnect ALL patient leads, test leads, and DUT outlet connections.
2. Press CAL to begin calibration, as shown:



If the calibration fails, the previously stored readings will be used until a passing calibration has occurred. Also, the esc/stop key has no effect during calibration.

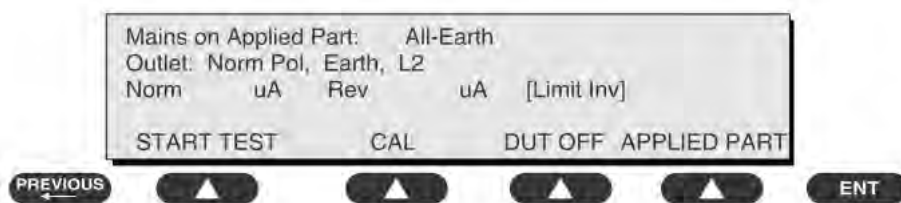
3. When the calibration is finished, the Mains on Applied Part test will reappear.

WARNING

- A 2-beep-per-second signal indicates high voltage present at the applied part terminals while a calibration is being performed.
 - High voltage is present at applied part terminals while measurements are being taken.
-

To Perform the Test

1. From the MAIN MENU, or with the outlet unpowered, plug the DUT into the 601
2. Attach the applied parts to the 601PRO applied part terminals.
3. Attach the red terminal lead to a conductive part on the DUT enclosure.
4. Press shortcut key 7. The Mains on Applied Part test is displayed.



5. Select the desired outlet configuration and applied part to test using the appropriate SOFT KEYS:
6. Press START TEST (SOFT KEY 1) to begin the test.
7. Press the print data key to generate a printout of the latest measurement.

NOTE

- **If all of the applied parts correspond to the instrument type, the applied parts will be tied together and one reading will be taken. If any of the applied parts differ from the instrument type, all applied parts will be tested individually, based on the type of applied part. This applies to Auto and Step modes only.**
-

In Case of Failure

- Check any broken of the enclosure. Replace any defective part.
- Inspect wiring for bad crimps, poor connections, or damage.
- Test the wall outlet; verify it is grounded and is free of other wiring abnormalities. Notify the user or owner to correct any deviations. As a work around, check the other outlets to see if they could be used instead.
- Change another probe to confirm if the fail is caused by console.
- If the leakage current measurement tests fail on a new unit and if situation can not be corrected, submit a Safety Failure Report to document the system problem. Remove unit from operation.
- If all else fails, stop using and inform the Customer Service Engineer for analysis and disposal.

LIMITS

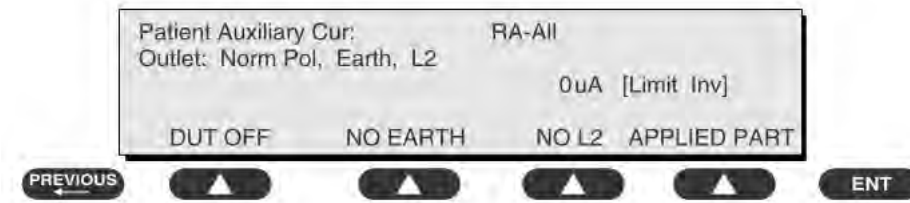
- For CF  applied parts: 50 μ A
- For BF  applied parts: 5000 μ A

A.8 Patient Auxiliary Current

Patient Auxiliary currents are measured between any selected ECG jack and the remaining selected ECG jacks. All measurements may have a true RMS only response.

Preparation

1. From the MAIN MENU, or with the outlet unpowered, plug the DUT into the 601PRO front panel outlet, and turn on the device.
2. Attach the patient leads to the 601PRO ECG jacks.
3. Define the Lead Types from the View Settings Option (refer to: Lead Type Definitions in Section 5 of this chapter).
4. Press shortcut key 8. The Patient Auxiliary Current test is displayed, and the test begins immediately. Display values are continuously updated until another test is selected.



5. Press SOFT KEYS 1-4 to select leakage tests
6. Press APPLIED PART (SOFT KEY 4) at any time to select the desired applied part leakage current:
7. Modify the configuration of the front panel outlet by pressing the appropriate SOFT KEY on the 601PRO:
8. Press the print data key at any time to generate a printout of the latest measurement.

In Case of Failure

- Check any broken of the enclosure. Replace any defective part.
- Inspect wiring for bad crimps, poor connections, or damage.
- Test the wall outlet; verify it is grounded and is free of other wiring abnormalities. Notify the user or owner to correct any deviations. As a work around, check the other outlets to see if they could be used instead.
- Change another probe to confirm if the fail is caused by console.
- If the leakage current measurement tests fail on a new unit and if situation can not be corrected, submit a Safety Failure Report to document the system problem. Remove unit from operation.
- If all else fails, stop using and inform the Customer Service Engineer for analysis and disposal.

LIMITS

For CF  applied parts,

- ◆ 10 μ A in Normal Condition
- ◆ 50 μ A in Single Fault Condition

For BF  applied parts,

- ◆ 100 μ A in Normal Condition
- ◆ 500 μ A in Single Fault Condition

A.9 Scheduled Electrical Safety Inspection

For scheduled electrical safety inspection, test items 1, 2, 3, 4, 5, 6, 7, and 8 included in the **ELECTRICAL SAFETY INSPECTION FORM** shall be performed.

ELECTRICAL SAFETY INSPECTION FORM

Location:				Technician:	
Equipment:				Control Number:	
Manufacturer:		Model:		SN:	
Measurement equipment /SN:				Date of Calibration:	
INSPECTION AND TESTING				Pass/Fail	Limit
1	Power Cord Plug				
2	Device Enclosure and Accessories				
3	Device Labeling				
4	Protective Earth Resistance		___ Ω		Max 0.2 Ω
5	Earth Leakage	Normal condition (NC)	___ μA		Max: NC: 300 μA (refer to UL60601-1) * NC: 500 μA (refer to IEC60601-1) * SFC: 1000 μA
		Single Fault condition (SFC)	___ μA		
6	Patient Leakage Current	Normal condition (NC)	<input type="checkbox"/> BF ___ μA		Max: CF applied part: NC:10 μA , SFC: 50 μA BF applied part: NC:100 μA , SFC: 500 μA
			<input type="checkbox"/> CF ___ μA		
		Single Fault condition (SFC)	<input type="checkbox"/> BF ___ μA		
			<input type="checkbox"/> CF ___ μA		
7	Mains on Applied Part Leakage		<input type="checkbox"/> BF ___ μA		Max: CF applied part: 50 μA BF applied part: 5000 μA
			<input type="checkbox"/> CF ___ μA		
8	Patient Auxiliary Current	Normal condition (NC)	<input type="checkbox"/> BF ___ μA		Max: CF applied part: NC:10 μA , SFC: 50 μA BF applied part: NC:100 μA , SFC: 500 μA
			<input type="checkbox"/> CF ___ μA		
		Single Fault condition (SFC)	<input type="checkbox"/> BF ___ μA		
			<input type="checkbox"/> CF ___ μA		

Note: The equipment sold to the United States shall comply with the requirement of UL60601-1; others shall comply with the requirement of IEC60601-1.

Name/ Signature: _____ Date: _____

A.10 Electrical Safety Inspection after Repair

The following table specifies test items to be performed after the equipment is repaired.

Repair with main unit not disassembled		Test items: 1, 2, 3
Repair with main unit disassembled	When neither power supply PCBA nor patient electrically-connected PCBA is repaired or replaced	Test items: 1, 2, 3, 4
	When power supply PCBA is repaired or replaced	Test items: 1, 2, 3, 4, 5
	When patient electrically-connected PCBA is repaired or replaced	Test items: 1, 2, 3, 4, 6, 7, 8
	When both power supply PCBA and patient electrically-connected PCBA are repaired or replaced	Test items: 1, 2, 3, 4, 5, 6, 7, 8

ELECTRICAL SAFETY INSPECTION FORM

Location:				Technician:	
Equipment:				Control Number:	
Manufacturer:		Model:		SN:	
Measurement equipment /SN:				Date of Calibration:	
INSPECTION AND TESTING				Pass/Fail	Limit
1	Power Cord Plug				
2	Device Enclosure and Accessories				
3	Device Labeling				
4	Protective Earth Resistance		____Ω		Max 0.2 Ω
5	Earth Leakage	Normal condition(NC)	____μA		Max: NC: 300μA(refer to UL60601-1) * NC: 500μA(refer to IEC60601-1) * SFC: 1000μA
		Single Fault condition(SFC)	____μA		
6	Patient Leakage Current	Normal condition(NC)	<input type="checkbox"/> BF ____μA		Max: CF applied part: NC:10μA, SFC: 50μA BF applied part: NC:100μA, SFC: 500μA
			<input type="checkbox"/> CF ____μA		
		Single Fault condition(SFC)	<input type="checkbox"/> BF ____μA		
			<input type="checkbox"/> CF ____μA		
7	Mains on Applied Part Leakage		<input type="checkbox"/> BF ____μA		Max: CF applied part: 50μA BF applied part: 5000μA
			<input type="checkbox"/> CF ____μA		
8	Patient Auxiliary Current	Normal condition(NC)	<input type="checkbox"/> BF ____μA		Max: CF applied part: NC:10μA, SFC: 50μA BF applied part: NC:100μA, SFC: 500μA
			<input type="checkbox"/> CF ____μA		
		Single Fault condition(SFC)	<input type="checkbox"/> BF ____μA		
			<input type="checkbox"/> CF ____μA		

Note: The equipment sold to the United States shall comply with the requirement of UL60601-1; others shall comply with the requirement of IEC60601-1.

Name/ Signature: _____ Date: _____

FOR YOUR NOTES

